



# Pneumococcal Disease, Invasive (IPD)

## CHAPTER CONTENT

1. [Case Definition](#)
2. [Diagnosis](#)
3. [Reporting](#)
4. [Overview](#)
5. [Public Health Measures](#)
6. [Education](#)
7. [Epidemiology](#)
8. [References](#)

The following chapter is adapted with permission from Alberta Health, for additional guidance related to the management of Pneumococcal disease Invasive see Alberta Public Health Disease Management Guidelines: [Pneumococcal Disease Invasive](#)

## 1. CASE DEFINITION

### Confirmed Case

- Clinical illness\* or evidence of invasive disease with laboratory confirmation of infection:
  - Isolation of *Streptococcus pneumoniae* from a normally sterile site\*\* (not including the middle ear and pleural cavity) **OR**
  - Detection of *S. pneumoniae* DNA by specific nucleic acid test (e.g., polymerase chain reaction [PCR]) from a normally sterile site (excluding the middle ear and pleural cavity)

### Probable Case

- Clinical illness\* or evidence of invasive disease with no other apparent cause and with non-confirmatory laboratory evidence:
  - Demonstration of *S. pneumoniae* antigen from a normally sterile site (excluding the middle ear).

\* Clinical illness associated with invasive pneumococcal disease manifests mainly as: pneumonia with bacteremia, bacteremia without a known site of infection, and meningitis.

\*\*Normally sterile site specimens are defined as:

- Blood
- Cerebrospinal fluid (CSF)
- Pleural fluid



- Peritoneal fluid
- Pericardial fluid
- Bone
- Joint fluid **OR**
- Specimens taken during surgery (e.g., muscle collected during debridement for necrotizing fasciitis or fluid from a deep abscess)

#### NOTE:

- A specimen taken from a non-sterile site collected during a sterile procedure is not considered a “normally sterile site”.
- Sputum and gastric or bronchial lavages are not considered sterile specimens.
- The middle ear is not considered a sterile site.

## 2. DIAGNOSIS

- A diagnosis of invasive pneumococcal disease is made by the isolation of *S. pneumoniae* from a normally sterile site excluding the middle ear.
- Recovery of pneumococci from an upper respiratory tract culture is not useful in patients with otitis media, pneumonia, or sinusitis.
- Blood cultures should be obtained, and cultures of other appropriate fluids (e.g., CSF, pleural fluid) may also be indicated.
- Sensitivity of strains should be determined by Alberta Provincial Lab.
- For more information, refer to the
  - [Alberta Public Health Laboratories Guide to Services](#)
  - [Government of Canada Guide to Services](#)
  - Alberta Public Health Disease Management Guidelines: [Pneumococcal Disease Invasive](#)

## 3. REPORTING

### Health Care Professionals

- Confirmed or probable cases are to be reported to the Office of the Chief Public Health Officer (OCPHO) by telephone (867) 920-8646, fax (867) 873-0442, or email **within 24 hours** after diagnosis is made, or opinion is formed **AND**
- Complete and fax (867) 873-0442 the [Communicable Disease Reporting Form](#) to the OCPHO within **24 hours**
- **Immediately** report all outbreaks or suspect outbreaks by telephone (867)-920-8646 to the OCPHO.



### Laboratories

- Report all positive results to the OCPHO by fax (867) 873-0442, **within 24 hours**

## 4. OVERVIEW

### Causative Agent

- *S. pneumoniae* is a gram-positive diplococcus of the *Streptococcaceae* family, and typically presents in pairs or short chains.
- There are around 90 serotypes.
- Most serotypes of *S. pneumoniae* can cause disease, but only a few produce the majority of invasive pneumococcal infection.
- Capsular polysaccharides make up the virulence factors of the bacteria.

### Clinical Presentation and Major Complications

- For information regarding Clinical Presentation and Major Complications see Alberta Public Health Disease Management Guidelines: [Pneumococcal Disease Invasive](#).

### Transmission

- Usual mode of transmission is person to person via droplets spread through coughing, sneezing, or direct oral contact.
- May also be transmitted indirectly via articles (fomites) contaminated with infected saliva or respiratory secretions.
- *S. pneumoniae* can survive up to 25 days in dust, 1-11 days on glass or similar impermeable surfaces, and seven days in sputum.
- Those at highest risk of acquiring serious invasive pneumococcal disease (IPD) are individuals who are living with:
  - Chronic medical conditions, including anatomic or functional asplenia,
  - Sickle cell disease
  - Chronic cardiovascular disease
  - Diabetes mellitus
  - Cirrhosis
  - Cancer and/or cancer treatment
  - Lymphomas, including Hodgkin's disease
  - Multiple myeloma
  - Chronic renal failure or nephrotic syndrome
  - HIV infection
  - Organ transplant
  - Other causes of immunosuppression
- Pneumococcal infections are most common during the winter and spring.
- *S. pneumoniae* is often found colonized in the upper respiratory tract of healthy people; these



carriers are the reservoir for these bacteria.

- It has been estimated that 40% of individuals become carriers of the bacteria by 1 year old.
- The duration of colonization varies but is generally longer in adults than children.
- Children who attend childcare centres have a higher carrier rate due to the increased frequency and level of contact with other children.
- Adults, who are in frequent contact with young children, are at higher risk of pneumococcal disease as children are more likely to be colonized with the bacteria.
- The prevalence of asymptomatic carriage varies with age and the incidence of upper respiratory infections.
- Infection generally confers immunity to the specific serotype, and this immunity may last for many years.
- The best way to assure immunity to multiple serotypes of *S. pneumoniae* is through immunization.

#### Incubation Period

- The incubation period may be as short as 1-3 days, depending on the type of infection.
- The period of communicability is variable but may persist as long as the organism is present in the respiratory tract.
- Individuals are no longer considered infectious 24 hours following initiation of appropriate antibiotics.

#### Clinical Guidance

- For patient-specific clinical management consult your local healthcare professional, paediatrician, infectious disease specialist, and/or the [NWT Clinical Practice Guidelines](#) (chapter 3 and chapter 8).

## 5. PUBLIC HEALTH MEASURES

#### Key Investigation

- Determine immunization status
- Obtain a history of illness including symptoms, and date of symptom onset
- Determine possible source of infection, taking into consideration the incubation period
- Identify underlying medical conditions that could predispose to invasive disease

#### Management of Cases

- Encourage proper hand hygiene and respiratory etiquette
- Practice Routine precautions for hospitalized individuals
- Droplet precautions may be warranted when antibiotic resistant infection is present.
- For cases who have not been previously vaccinated the following is recommended
  - 1 dose of Pneu-C-13 AND
  - 1 dose of Pneu-P-23 **at least 8 weeks** after Pneu-C-13 vaccine Immunization



### Management of Contacts

- Transmission through casual contact is unlikely
- No specific contact management is recommended
- Immunization for Outbreak Control
  - During outbreaks of pneumococcal infection due to Pneu-C-13 vaccine serotypes, immunization with Pneu-C-13 vaccine is recommended for children who have not been adequately immunized with Pneu-C-13 vaccine.
  - Pneu-P-23 or Pneu-C-13 vaccine can be used in adults, if the outbreak is due to serotypes included in the vaccine.

### Prevention

- Pneumococcal disease is vaccine preventable.
- Certain risk factors or chronic conditions put children and adults at an increased risk of acquiring invasive pneumococcal disease.
- Individuals with increased risk factors should be assessed for eligibility according to the [Evergreen Canadian Immunization Guide](#).
- Vaccines available for prevention of disease include pneumococcal polysaccharide 23-valent vaccine (Pneu-P-23) and pneumococcal conjugate 13-valent vaccine (Pneu-C-13).
- The NWT provides publicly funded immunization programs for pneumococcal disease.
- For more information on NWT pneumococcal vaccine program please follow the [NWT Immunization Schedule](#).

## 6. PUBLIC & HEALTH PROFESSIONAL EDUCATION

For more information about invasive pneumococcal disease:

- Health Canada: [Canada/invasive pneumococcal disease](#)
- Centers For Disease Control and Prevention: [CDC/invasive pneumococcal disease](#)
- Alberta Public Health Disease Management Guidelines: [Pneumococcal Disease Invasive](#)

## 7. EPIDEMIOLOGY

- For information on the epidemiology of invasive pneumococcal disease in the Northwest Territories (NWT) see: [Epidemiological Summary of Communicable Diseases HSS Professionals](#)

## 8. REFERENCES

The information from this chapter is adapted, with permission, from Alberta Health. For more information regarding Invasive Pneumococcal Disease see: [Alberta Public Health Disease Management Guidelines: Pneumococcal Disease, Invasive](#)

1. Canadian Immunization Guide:



- <https://www.canada.ca/en/public-health/services/canadian-immunization-guide.html>
2. Government of Canada- Invasive Pneumococcal Disease for Health Professionals:  
<https://www.canada.ca/en/public-health/services/immunization/vaccine-preventable-diseases/invasive-pneumococcal-disease/health-professionals.html>