

**HIV/AIDS Case Report**  
**Adult, Adolescent and Pediatric (non maternal-fetal) Cases**  
(A separate form is available for maternal-fetal transmission cases.)

**GENERAL INFORMATION**

- The Canadian AIDS Case Reporting Surveillance System is a voluntary collaboration of all provinces and territories. AIDS reports are collected by the provinces and territories under the authority of each provincial or territorial Health Act. The system is coordinated by the Division of HIV/AIDS Surveillance at the Bureau of HIV/AIDS and STD, Laboratory Centre for Disease Control, Health Canada.

**INSTRUCTIONS**

- Pediatric cases include children who are less than 15 years old at time of diagnosis of HIV or AIDS, whichever is being reported.
- Complete a new form at each change of status eg. HIV to AIDS, death update, or new disease information. If you are uncertain whether a report form has been previously submitted, regardless of source, please submit a new report form.
- PLEASE PRESS HARD AND USE A BALL POINT PEN
- Physicians in Quebec:** After completing the HIV/AIDS Case Report form, please forward to the Local Medical Officer of Health.  
**Physicians in all other provinces/territories:** After completing the HIV/AIDS Case Report form, please forward copies 2,3 and 4 to the Local Medical Officer of Health.
- Cases of AIDS must meet the Canadian surveillance case definition as described in the: [Canada Diseases Weekly Report 1987;13-38:169-176](#) and the [Canada Communicable Disease Report 1993; 19-15:116-117](#).
- PLEASE DO NOT SEND CASE REPORTS DIRECTLY TO THE LABORATORY CENTRE FOR DISEASE CONTROL. ALL CASES MUST BE REPORTED TO THE LOCAL MEDICAL OFFICERS OF HEALTH.**

**SECTION I - PATIENT INFORMATION**

- 1 Death Update:**
- If this form is being used as a death update, please provide initials, date of birth and date of death.
  - Use Section IV to indicate all diseases present at death.
  - If a complete case report form has been previously submitted and there is no new information, it is not necessary to complete Sections II, III, V, and VI.

**SECTION III - LABORATORY DATA**

- Laboratory evidence of HIV infection in persons over 15 months of age or in non maternal-fetal transmission**  
For the purpose of surveillance, a confirmed, repeatedly reactive screening test for HIV antibody constitutes sufficient laboratory evidence of HIV infection in any person over 15 months of age or in infants less than 15 months of age if maternal-fetal transmission is not suspected. Other acceptable evidence is outlined in the [Canada Diseases Weekly Report 1987;13-38:169-176](#) and the [Canada Communicable Disease Report 1993; 19-15:116-117](#).
- In the absence of laboratory evidence for HIV infection, causes of immunodeficiency that DISQUALIFY diseases as indicators of AIDS are:**
  - High-dose or long-term systemic corticosteroid therapy or other immunosuppressive/cytotoxic therapy within 3 months before the onset of the indicator disease.
  - Any of the following diseases diagnosed <3 months after diagnosis of the indicator disease: Hodgkin's disease, non-Hodgkin's lymphoma (other than primary brain lymphoma), lymphocytic leukemia, multiple myeloma, any other cancer of lymphoreticular tissue, or angioimmunoblastic lymphadenopathy.
  - A genetic (congenital) immunodeficiency syndrome or an acquired immunodeficiency syndrome atypical of HIV infection, such as one involving hypogammaglobulinemia.

**SECTION IV - DISEASES INDICATIVE OF AIDS**

- The following definitions are intended for purposes of surveillance. They are not intended to provide clinical guidance in diagnosis and should not be used in that way.
- For classification under the surveillance definition for AIDS, most diseases may be diagnosed by presumptive methods if the patient **has had a positive HIV test**.
- For Encephalopathy (HIV-related) and Wasting syndrome due to HIV, the methods of diagnosis described here are not truly definitive, but are sufficiently rigorous for surveillance purposes.

<p><b>Bacterial pneumonia, recurrent</b></p> <p><b>Definitive</b></p> <ul style="list-style-type: none"> <li>Recurrent (more than one episode in a 1-year period), acute (new X-ray evidence, not present earlier) pneumonia diagnosed by both of the following:           <ol style="list-style-type: none"> <li>culture (or other organism-specific diagnostic method) obtained from a clinically reliable specimen of a pathogen that typically causes pneumonia (other than <i>Pneumocystis carinii</i> or <i>Mycobacterium tuberculosis</i>); and</li> <li>radiologic evidence of pneumonia. Cases that do not have laboratory confirmation of a causative organism for one of the episodes of pneumonia will be considered to be presumptively diagnosed.</li> </ol> </li> </ul> <p><b>Presumptive</b></p> <ul style="list-style-type: none"> <li>Recurrent (more than one episode in a 1-year period), acute (new symptoms, signs, or X-ray evidence not present earlier) pneumonia diagnosed on clinical grounds by the patient's physician.</li> </ul>	<p><b>Isosporiasis, chronic intestinal (&gt;1 mo. duration)</b></p> <p><b>Definitive</b></p> <ul style="list-style-type: none"> <li>Microscopy (histology or cytology).</li> </ul> <p><b>Kaposi's sarcoma</b></p> <p><b>Definitive</b></p> <ul style="list-style-type: none"> <li>Microscopy (histology or cytology).</li> <li>Characteristic gross appearance of an erythematous or violaceous plaque-like lesion on skin or mucous membrane. <b>(Note: presumptive diagnosis of Kaposi's sarcoma should not be made by clinicians who have seen few cases of it.)</b></li> </ul> <p><b>Lymphoma, Burkitt's (or equivalent term)</b> <b>Lymphoma, immunoblastic (or equivalent term)</b> <b>Lymphoma, primary in brain</b></p> <p><b>Definitive</b></p> <ul style="list-style-type: none"> <li>Microscopy (histology or cytology).</li> </ul> <p><b>Mycobacterium avium complex or M. kansasii (disseminated or extrapulmonary)</b> <b>Mycobacterium of other species or unidentified species</b></p> <p><b>Definitive</b></p> <ul style="list-style-type: none"> <li>Culture</li> </ul> <p><b>Presumptive</b></p> <ul style="list-style-type: none"> <li>Microscopy of a specimen from stool or normally sterile body fluids or tissue from a site other than lungs, skin, or cervical or hilar lymph nodes, showing acid-fast bacilli of a species not identified by culture.</li> </ul> <p><b>M. tuberculosis (disseminated or extrapulmonary)</b></p> <p><b>Definitive</b></p> <ul style="list-style-type: none"> <li>Culture</li> </ul> <p><b>M. tuberculosis (pulmonary)</b></p> <p><b>Definitive</b></p> <ul style="list-style-type: none"> <li>Culture</li> </ul> <p><b>Presumptive</b></p> <ul style="list-style-type: none"> <li>When bacteriologic confirmation is not available, other reports may be considered to be verified cases of pulmonary tuberculosis if the criteria meet those in the definitions in any of the following sources:           <ol style="list-style-type: none"> <li>the Canada Diseases Weekly Report 1991;17S3:32-3;</li> <li>the 1990 Canadian Tuberculosis Reporting System form; or</li> <li>Health Reports 1992;4(2) (Suppl.No10).xi.</li> </ol> </li> </ul> <p><b>Pneumocystis carinii pneumonia</b></p> <p><b>Definitive</b></p> <ul style="list-style-type: none"> <li>Microscopy (histology or cytology).</li> </ul> <p><b>Presumptive</b></p> <ul style="list-style-type: none"> <li>A history of dyspnea on exertion or nonproductive cough of recent onset (within the past 3 months); and chest X-ray evidence of diffuse bilateral interstitial infiltrates or gallium scan evidence of diffuse bilateral pulmonary disease; and arterial blood gas analysis showing an arterial pO<sub>2</sub> &lt; 70mm Hg or a low respiratory diffusing capacity (&lt; 80% of predicted values) or an increase in the alveolar-arterial oxygen tension gradient; and no evidence of bacterial pneumonia.</li> </ul>
<p><b>Candidiasis (bronchi, trachea or lungs)</b></p> <p><b>Definitive</b></p> <ul style="list-style-type: none"> <li>Gross inspection by endoscopy, autopsy or microscopy (histology or cytology) on a specimen obtained directly from the tissues affected (including scrapings from the mucosal surface), not from a culture.</li> </ul> <p><b>Candidiasis (esophageal)</b></p> <p><b>Definitive</b></p> <ul style="list-style-type: none"> <li>Gross inspection by endoscopy, autopsy or microscopy (histology or cytology) on a specimen obtained directly from the tissues affected (including scrapings from the mucosal surface), not from a culture.</li> </ul> <p><b>Presumptive</b></p> <ul style="list-style-type: none"> <li>Recent onset of retrosternal pain on swallowing; and oral candidiasis diagnosed by the gross appearance of white plaques on an erythematous base or by the microscopic appearance of fungal mycelial filaments in an uncultured specimen scraped from the oral mucosa.</li> </ul>	<p><b>Progressive multifocal leukoencephalopathy</b></p> <p><b>Definitive</b></p> <ul style="list-style-type: none"> <li>Microscopy (histology or cytology).</li> </ul> <p><b>Salmonella septicemia, recurrent</b></p> <p><b>Definitive</b></p> <ul style="list-style-type: none"> <li>Culture</li> </ul> <p><b>Toxoplasmosis of brain</b></p> <p><b>Definitive</b></p> <ul style="list-style-type: none"> <li>Microscopy (histology or cytology).</li> </ul> <p><b>Presumptive</b></p> <ul style="list-style-type: none"> <li>Recent onset of a focal neurologic abnormality consistent with intracranial disease or a reduced level of consciousness; and brain imaging evidence of a lesion having a mass effect (on computed tomography or nuclear magnetic resonance) or the radiographic appearance of which is enhanced by injection of contrast medium; and serum antibody to toxoplasmosis or successful response to therapy for toxoplasmosis.</li> </ul>
<p><b>Cervical cancer, invasive</b></p> <p><b>Definitive</b></p> <ul style="list-style-type: none"> <li>Microscopy (histology or cytology).</li> </ul> <p><b>Coccidioidomycosis (disseminated or extrapulmonary)</b> <b>Cryptococcosis (extrapulmonary)</b></p> <p><b>Definitive</b></p> <ul style="list-style-type: none"> <li>Microscopy (histology or cytology), culture or detection of antigen in a specimen obtained directly from the tissues affected or a fluid from those tissues.</li> </ul> <p><b>Cryptosporidiosis (chronic intestinal, &gt;1 mo. duration)</b></p> <p><b>Definitive</b></p> <ul style="list-style-type: none"> <li>Microscopy (histology or cytology).</li> </ul> <p><b>Cytomegalovirus disease (other than in liver, spleen or nodes)</b></p> <p><b>Definitive</b></p> <ul style="list-style-type: none"> <li>Microscopy (histology or cytology), culture or detection of antigen in a specimen obtained directly from the tissues affected or a fluid from those tissues.</li> </ul> <p><b>Cytomegalovirus retinitis (with loss of vision)</b></p> <p><b>Definitive</b></p> <ul style="list-style-type: none"> <li>Microscopy (histology or cytology), culture or detection of antigen in a specimen obtained directly from the tissues affected or a fluid from those tissues.</li> </ul> <p><b>Presumptive</b></p> <ul style="list-style-type: none"> <li>Characteristic appearance on serial ophthalmoscopic examinations (e.g., discrete patches of retinal whitening with distinct borders, spreading in a centrifugal manner following blood vessels, progressing over several months, frequently associated with retinal vasculitis, hemorrhage, and necrosis). Resolution of active disease leaves retinal scarring and atrophy with retinal pigment epithelial mottling.</li> </ul>	<p><b>Wasting syndrome due to HIV</b></p> <p><b>Definitive</b></p> <ul style="list-style-type: none"> <li>Findings of profound involuntary weight loss (more than 10% of baseline body weight) plus either chronic diarrhea (2 or more loose stools per day for 30 days or more) or chronic weakness and documented fever (30 days or more, intermittent or constant) in the absence of concurrent illness or condition other than HIV infection that could explain the findings (e.g., cancer, tuberculosis, cryptosporidiosis, or other specific enteritis).</li> </ul>
<p><b>Encephalopathy, HIV-related (dementia)</b></p> <p><b>Definitive</b></p> <ul style="list-style-type: none"> <li>Clinical findings of disabling cognitive and/or motor dysfunction interfering with occupation or activities of daily living in an adult, or the loss of behavioural development milestones affecting a child, progressing over weeks to months in the absence of a concurrent illness or condition other than HIV infection that could explain the findings. Methods to rule out such concurrent illnesses and conditions must include cerebrospinal fluid examination and either brain imaging (computed tomography or magnetic resonance) or autopsy.</li> </ul> <p><b>Herpes simplex: chronic ulcer(s) (&gt;1 mo. duration) or bronchitis, pneumonitis or esophagitis</b> <b>Histoplasmosis (disseminated or extrapulmonary)</b></p> <p><b>Definitive</b></p> <ul style="list-style-type: none"> <li>Microscopy (histology or cytology), culture or detection of antigen in a specimen obtained directly from the tissues affected or a fluid from those tissues.</li> </ul>	
<b>Diseases affecting pediatric cases only (&lt;15 years old)</b>	
<p><b>Bacterial infections, multiple or recurrent (excluding recurrent bacterial pneumonia)</b></p> <p><b>Definitive</b></p> <ul style="list-style-type: none"> <li>Culture</li> </ul>	<p><b>Lymphoid interstitial pneumonia and/or Pulmonary lymphoid hyperplasia</b></p> <p><b>Definitive</b></p> <ul style="list-style-type: none"> <li>Microscopy (histology or cytology).</li> </ul> <p><b>Presumptive</b></p> <ul style="list-style-type: none"> <li>Bilateral reticulonodular interstitial pulmonary infiltrates present on chest X-ray ≥ 2 months with no response to antibiotic treatment.</li> </ul>



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Protected when completed

### HIV/AIDS Case Report Adult, Adolescent and Pediatric (non maternal-fetal) Cases

HIV     AIDS     New case report     Update

For provincial/territorial use	For use by LCDC
Provincial ID Number	EPIC No.
Province to which case is attributed	Date received YY MM DD

#### SECTION I - PATIENT INFORMATION

Reporting physician's name		City	Telephone number ( )
Hospital or clinic		City	Province
Is another physician providing ongoing care to this patient? Name		If so, please provide name, city and telephone number. City Telephone number ( )	
Patient's initials First Middle Last	Sex <input type="checkbox"/> M <input type="checkbox"/> F	Date of birth YY MM DD	Vital Status <input type="checkbox"/> Alive (If yes, date last known to be alive) <input type="checkbox"/> Dead (If yes, date of death) <input type="checkbox"/> Unknown
<p>• Is the patient: (please ask patient to assist you in answering this question)</p> <input type="checkbox"/> White <input type="checkbox"/> South Asian (e.g. East Indian, Pakistani, Sri Lankan, Punjabi, Bangladeshi, etc.) <input type="checkbox"/> Black (e.g. African, Haitian, Jamaican, Somali, etc.) <input type="checkbox"/> Arab/West Asian (e.g., Armenian, Egyptian, Iranian, Lebanese, Moroccan, etc.) <input type="checkbox"/> North American Indian <input type="checkbox"/> Métis <input type="checkbox"/> Inuit <input type="checkbox"/> Latin-American (e.g. Mexican, Central/South American, etc.) <input type="checkbox"/> Asian (e.g., Chinese, Japanese, Vietnamese, Cambodian, Indonesian, Laotian, Korean, Filipino, etc.) <input type="checkbox"/> Other - includes mixed ethnicity (specify) →			
What language does this person speak most often at home?		Country of birth <input type="checkbox"/> Canada <input type="checkbox"/> Other (specify) →	Year of arrival in Canada
City and province/territory of residence at diagnosis City Province/Territory First 3 digits of Postal Code		Current city and province/territory of residence City Province/Territory First 3 digits of Postal Code	

#### SECTION II - RISK(S) ASSOCIATED WITH THE TRANSMISSION OF HIV IN THIS PATIENT

• Since January 1978 and preceding the diagnosis of HIV/AIDS, this patient had: (check ALL that apply)

Yes	No	Unknown	
<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	Sex with a male.
<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	Sex with a female.
<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<b>Heterosexual</b> sex with: (check ALL that apply)
<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	• an injection drug user;
<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	• a bisexual male;
<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	• a transfusion recipient with documented HIV infection;
<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	• a person with hemophilia/coagulation disorder;
<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	• a person born in a country where heterosexual transmission predominates. If yes, specify country;
<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	• a person with confirmed or suspected HIV infection or AIDS (whether or not risk factor is known).
<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	Injected non-prescription drugs (including steroids).
<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	Received pooled concentrates of factor VIII or IX for treatment of hemophilia/coagulation disorder. If yes, please complete Section 1 of the Supplement to HIV/AIDS Case Report.
<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	Received transfusion of whole blood or blood components such as packed red cells, plasma, platelets or cryoprecipitate. If yes, please complete Section 2 of the Supplement to HIV/AIDS Case Report.
<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	Exposure to HIV-contaminated blood or body fluids or concentrated virus in an occupational setting. If yes, specify occupation. →
<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	Other medical exposure (eg: organ or tissue transplant, artificial insemination). If yes, please give details in Section VI "Additional Information or Comments".
<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	Non-medical, non-occupational exposure which could have been the source of the infection (eg: acupuncture, tattoo, body piercing, breast milk). If yes, please give details of type of exposure, date and location in Section VI "Additional Information or Comments".

Since January 1978, has this patient donated blood, plasma, platelets, organs, tissues, semen or breast milk?  Yes  No  Unknown  
If yes, please give details of type of donation, date and location in Section VI "Additional Information or Comments".

Has the Red Cross or other appropriate donor program been notified?  Yes  No  Unknown

Do you want a public health official to ensure this notification?  Yes  No  Unknown

HC/SC 4205 E (08-96)

Distribution: White - Medical Officer of Health    Yellow - Ministry of Health    Pink - LCDC

**SECTION III - LABORATORY DATA**

• Does this case have evidence, as defined in the above instructions, of HIV infection?  
 Yes  No  Unknown

Date of first positive HIV test (if known)  
 Year Month

Current CD4 count (if known)  
 cells/ $\mu$ l

**SECTION IV - DISEASES INDICATIVE OF AIDS**

DISEASES	Date of Diagnosis		Diagnostic method		DISEASES	Date of Diagnosis		Diagnostic method	
	Year	Month	Definitive	Presumptive		Year	Month	Definitive	Presumptive
Bacterial pneumonia, recurrent			<input type="checkbox"/>	<input type="checkbox"/>	<i>Mycobacterium avium</i> complex or <i>M. kansasii</i> (disseminated or extrapulmonary)			<input type="checkbox"/>	<input type="checkbox"/>
Candidiasis (bronchi, trachea or lungs)			<input type="checkbox"/>	<input type="checkbox"/>	Mycobacterium of other species or unidentified species			<input type="checkbox"/>	<input type="checkbox"/>
Candidiasis (esophageal)			<input type="checkbox"/>	<input type="checkbox"/>	<i>M. tuberculosis</i> (disseminated or extrapulmonary) (Please complete SECTION V)			<input type="checkbox"/>	<input type="checkbox"/>
Cervical cancer, invasive			<input type="checkbox"/>	<input type="checkbox"/>	Specify Site: <input type="checkbox"/> Miliary <input type="checkbox"/> Pleurisy <input type="checkbox"/> Other respiratory <input type="checkbox"/> C.N.S. <input type="checkbox"/> Bone and joint <input type="checkbox"/> Genitourinary				
Coccidioidomycosis (disseminated or extrapulmonary)			<input type="checkbox"/>	<input type="checkbox"/>	Other (specify) →				
Cryptococcosis (extrapulmonary)			<input type="checkbox"/>	<input type="checkbox"/>	<i>M. tuberculosis</i> (pulmonary) (Please complete SECTION V)			<input type="checkbox"/>	<input type="checkbox"/>
Cryptosporidiosis (chronic intestinal, >1 mo. duration)			<input type="checkbox"/>	<input type="checkbox"/>	<i>Pneumocystis carinii</i> pneumonia			<input type="checkbox"/>	<input type="checkbox"/>
Cytomegalovirus disease (other than in liver, spleen or nodes)			<input type="checkbox"/>	<input type="checkbox"/>	Progressive multifocal leukoencephalopathy			<input type="checkbox"/>	<input type="checkbox"/>
Cytomegalovirus retinitis (with loss of vision)			<input type="checkbox"/>	<input type="checkbox"/>	Salmonella septicemia, recurrent			<input type="checkbox"/>	<input type="checkbox"/>
Encephalopathy, HIV-related (dementia)			<input type="checkbox"/>	<input type="checkbox"/>	Toxoplasmosis of brain			<input type="checkbox"/>	<input type="checkbox"/>
Herpes simplex: chronic ulcer(s) (>1 mo. duration) or bronchitis, pneumonitis or esophagitis			<input type="checkbox"/>	<input type="checkbox"/>	Wasting syndrome due to HIV			<input type="checkbox"/>	<input type="checkbox"/>
Histoplasmosis (disseminated or extrapulmonary)			<input type="checkbox"/>	<input type="checkbox"/>	<b>Diseases affecting pediatric cases only (&lt;15 years old)</b>				
Isosporiasis, chronic intestinal (>1 mo. duration)			<input type="checkbox"/>	<input type="checkbox"/>	Bacterial infections, multiple or recurrent (excluding recurrent bacterial pneumonia)			<input type="checkbox"/>	<input type="checkbox"/>
Kaposi's sarcoma			<input type="checkbox"/>	<input type="checkbox"/>	Lymphoid interstitial pneumonia and/or Pulmonary lymphoid hyperplasia			<input type="checkbox"/>	<input type="checkbox"/>
Lymphoma, Burkitt's (or equivalent term)			<input type="checkbox"/>	<input type="checkbox"/>					
Lymphoma, immunoblastic (or equivalent term)			<input type="checkbox"/>	<input type="checkbox"/>					
Lymphoma, primary in brain			<input type="checkbox"/>	<input type="checkbox"/>					

**SECTION V - TUBERCULOSIS**

- Before the diagnosis of AIDS, was this patient ever treated for tuberculosis?  Yes - When? → Year Month  No  Unknown
- Has this patient ever had a PPD skin test?  Yes - What was the size in mm → mm  No  Unknown
- If the PPD test was negative, was the patient energy tested?  Yes  No  Unknown If yes, were any sites positive?  Yes  No  Unknown

**SECTION VI - ADDITIONAL INFORMATION OR COMMENTS**

(Please use this section for information of interest about the acquisition of the virus, etc.)

Person completing this form	Telephone number	Date report completed
	( )	YY MM DD

**FOR PROVINCIAL/TERRITORIAL USE: To which exposure category has this patient been assigned?**

- |                                                          |                                                    |                                                |                                                         |                                             |
|----------------------------------------------------------|----------------------------------------------------|------------------------------------------------|---------------------------------------------------------|---------------------------------------------|
| <input type="checkbox"/> Men who have sex with men (MSM) | <input type="checkbox"/> Injection drug user (IDU) | <input type="checkbox"/> MSM and IDU           | <input type="checkbox"/> Heterosexual - Endemic         | <input type="checkbox"/> NIR - Heterosexual |
| <input type="checkbox"/> Blood transfusion recipient     | <input type="checkbox"/> Clotting factor recipient | <input type="checkbox"/> Occupational exposure | <input type="checkbox"/> Heterosexual - Partner at risk | <input type="checkbox"/> NIR - Other        |