# **Notification Timeline:**

From Lab/Practitioner to Public Health: Within 48 hours. From Public Health to Saskatchewan Health: Within 2 weeks. Public Health Follow-up Timeline: Within 72 hours.

## Public Health Purpose for Notification of HIV

- To support positive outcomes for individuals and the community through:
  - Engagement in care, education about prevention and control measures, referrals to harm reduction services, and other communicable disease services including TB screening and immunizations;
- To identify cases of HIV through contact tracing in order to prevent further transmission;
- To offer testing and referral to supportive services to at risk individuals through contact tracing;
- To track epidemiology trends of HIV in Saskatchewan including risk factors and distribution;
- To identify locations where increased transmission of HIV may be occurring in order to inform other interventions;
- To monitor the effectiveness of prevention and control measures;
- To make timely and evidence informed actions on outbreaks; and
- To inform the public and medical community about HIV.

# **Surveillance Case Definition**<sup>1</sup> (Adapted from Public Health Agency of Canada, May 2008)

Confirmed Case:	Detection of HIV antibody with confirmation (e.g., EIA screening with		
Adults, Adolescents	confirmation by Western blot or other confirmatory test)		
and Children ≥ 18	OR		
months	detection of HIV nucleic acid (e.g., DNA PCR or plasma RNA)		
	OR		
	HIV p24 antigen with confirmation by neutralization assay		
	OR		

<sup>&</sup>lt;sup>1</sup> Surveillance case definitions ensure uniform reporting to allow comparability of surveillance data. The definition is not intended to be used for clinical or laboratory diagnosis or management of cases.

	isolation of HIV in culture.				
Confirmed Case:	Detection of HIV nucleic acid (e.g., DNA PCR or plasma RNA)				
Children < 18 OR					
months (on two	HIV p24 antigen with confirmation by neutralization assay				
separate samples	OR				
collected at different	isolation of HIV in culture.				
times)					
Probable Case:	Positive screening test that cannot be confirmed				
Adults	OR				
	indeterminate confirmatory test (HIV 1/2 Confirmatory assay or				
	Western blot)*				
	OR				
	reactive point of care test.				
Probable Case:	One positive confirmatory test without a second confirmatory test				
< 18 months	result available for the individual.				
In children < 18 months o	f age born to HIV-positive women, nucleic acid testing should be done within				
two weeks after birth and, if negative, repeated at 1 to 2 months and at 3 to 4 months of age. Any					
positive results should be repeated with a second specimen for confirmation.					
For children who are born to HIV-positive women and who have negative nucleic acid results, antibody					
testing should be done at 12 and 18 months of age to ensure that they have lost maternally derived					
antibodies. (This is not used to determine uninfected status but rather to eliminate the possibility of a					
	positive antibody result being misinterpreted.) These children should continue to be monitored until				
they have a negative HIV	antibody test.				

\*Indeterminate Western blot tests results on a repeat basis (3) are considered to be negative (U.S. Centers for Disease Control and Prevention, 1989).

from BC Center of Excellence in HIV [2018] and Vajpayee [2005])					
Stage	Criteria	CD4 at Diagnosis	AIDS-defining Illness		
0	Laboratory criteria met for acute HIV infection, or previous negative or				
	indeterminate HIV test within 180 days of first confirmed positive				
1	CD4≥500 AND No AIDS case report				
2	Stage 0 not met CD4 200-499				
3	AND CD4 <200 OR AIDS case report				

Table 1: Stage of HIV Infection at Diagnosis for individuals > 5 years of age (adapted from BC Center of Excellence in HIV [2018] and Vajpayee [2005])

One of the objectives is to identify individuals early in the course of infection to reduce further transmission to others. The CD4 count can be a marker to reflect stage of HIV infection at diagnosis.

No CD4 available

AND No AIDS case report

Unknown

# **Epidemiology and Occurrence**

Under development

# Additional Background Information

## **Causative Agent**

Human immunodeficiency virus. A retrovirus. Type 1 predominant in Canada, but Type 2 is present.

## **Reservoir/Source**

 Table 2: Fluids and tissues capable of transmitting blood borne pathogens (U.S. Centers for Disease Control, 2001)

Fluid	HIV
Blood and fluids visibly contaminated with blood	Yes
Semen	Yes
Vaginal secretions	Yes
Pleural, amniotic, pericardial, peritoneal, synovial and	Yes
cerebrospinal fluids and inflammatory exudates	
Saliva, faeces, nasal secretions, sputum, sweat, tears, urine,	No, unless contaminated
vomitus	with blood
Transplanted tissue or organs	Yes
Breast milk	Yes

# Symptoms

Individuals infected with HIV may experience several stages (Public Health Agency of Canada, 2013). The stage based on CD4 count (**Table 1**) is considered a more objective way to document stage. Below is a description of clinical presentation HIV based on stage of infection:

# • HIV Primary/Acute infection

Up to 90% of individuals experience symptoms within 2-4 weeks after infection (acute retroviral syndrome). Symptoms typically last 1-2 weeks but may last up to several months. These signs and symptoms include:

- fever (mean temperature 39.4°C [102.9°F] > 80%);
- arthralgia or myalgia, rash, lymphadenopathy, sore throat, fatigue, headache (40-80%);
- oral ulcers and/or genital ulcers, > 5 kg weight loss, nausea, vomiting, or diarrhea (10-40%).



# • Chronic Asymptomatic HIV infection

Many persons with HIV fall into this stage. It is the stage where the immune response is able to control viral replication and plasma viremia. In this stage of infection, people can experience the following signs and symptoms:

- generalized lymphadenopathy;
- thrombocytopenia.

# • Chronic Symptomatic HIV infection

This is the stage of profound immunosuppression. Signs and symptoms include:

- oral hairy leukoplakia;
- unexplained fever (> 2 weeks);
- fatigue or lethargy;
- unexplained weight loss (> 10% body weight);
- chronic diarrhea (> 3 weeks);
- unexplained lymphadenopathy (usually generalized);
- cervical dysplasia;
- dyspnea and dry cough;
- loss of vision;
- recurrent or chronic mucocutaneous candidiasis (oral, esophageal, vaginal);
- dysphagia (esophageal candidiasis);
- red/purple nodular skin or mucosal lesions (Kaposi sarcoma);
- encephalopathy;
- herpes zoster, especially if severe, multidermatomal or disseminated;
- increased frequency or severity of mucocutaneous herpes simplex virus infection;
- unexplained "anemia of chronic disease."

## Complications

Acquired immunodeficiency syndrome (AIDS). See Section 6-15.

## **Incubation Period**

The incubation period varies on each individual's ability to develop antibodies to HIV. Up to 90% of individuals experience symptoms within 2-4 weeks after infection. See <u>Symptoms.</u>

In HIV/AIDS research, the seroconversion period refers to the period of time it usually takes to develop detectable antibodies to HIV following infection with HIV. In 75% of persons, antibodies are produced in 4 to 8 weeks; in almost all persons, antibodies are produced within 14 weeks.



The seroconversion period is frequently described as the "window period." It is very significant in relation to the timing of HIV tests. In HIV testing, the window period refers to the time between a person becoming infected and when laboratory tests can detect HIV infection. The window period varies based on the test that is completed; progress in HIV testing technologies continues to result in tests with shorter window periods (British Columbia Centre for Disease Control, October, 2016).

Persons who are tested during the window period may receive a negative HIV test result although they may be infected with HIV. Persons disclosing HIV-related risk factors in the 14 weeks before testing negative for HIV are encouraged to be retested at the end of the window period.

In addition to test results, the risks that the individual has engaged in during the window period should be considered. Statistically it is very unlikely that a person with HIV would be tested during the 3 month window period (and test negative) however that possibility should be considered in persons with ongoing risk factors.

A summary of window periods based on the test used provides context to the reliability of the test results:

- antibody/antigen (4<sup>th</sup> generation test) has window period of approximately 2-3weeks;
- antibody test (3<sup>rd</sup> generation) has a window period of approximately 3-4 weeks;
- POCT has a window period of approximately 3-4 weeks;
- the Western blot or other confirmatory tests have a window period of approximately 4-6 weeks though it may take up to 8 weeks for a positive result.

Because window periods vary with the test, a negative test result at 3 months in an individual with no ongoing risk factors is deemed to be negative and no further testing is required.

# Period of Communicability

Communicability begins early after infection and extends throughout the individuals lifespan. Infectiousness is related to an individuals HIV viral load (i.e., high viral load increases potential for transmission). Generally, people are most infectious early and late in the course of infection. If the viral load is suppressed (<200 copies/mL), the risk of transmission is decreased. The presence if an STI does not increase the possibility of transmission if the HIV positive person is on effective ARVs (Barré-Sinoussi, 2018).

#### Mode of Transmission (Public Health Agency of Canada, 2010)

Transmission of HIV infection occurs essentially through specific exposure to blood or body fluids from an HIV-infected person. The risk of transmission decreases when the infected person is effectively responding to treatment.

In order to be infected, the virus must have an entry point, most directly through a person's bloodstream or mucous membranes (HIV cannot survive outside the body). HIV is transmitted from one person to another through:

- unprotected sexual intercourse (vaginal, anal or oral);
- shared needles, syringes or other equipment used for injecting drugs;
- unsterilized needles or equipment for tattooing, skin piercing or acupuncture;
- pregnancy, delivery and breast feeding (i.e., from an HIV-infected mother to her infant);
- occupational exposures in health care or other high risk settings.

# Table 4 Estimated Per-Act Probability of Acquiring HIV from a Known HIV-Infected Source by Exposure Act

Type of Exposure	Estimated Risk	Reference				
Parenteral						
Blood Transfusion	90% (9 in 10)					
Needle-sharing during injection drug use	0.63% (63 in 10000)	Patel, et al (2014)				
Percutaneous (needlestick)	0.23% (23 in 10 000)					
	Sexual					
Receptive anal intercourse	1.4% (7 in 5000)	Patel, et al (2014)				
Receptive penile-vaginal intercourse	0.08% (8 in 10000)	Patel, et al (2014)				
Insertive anal intercourse	0.11% (11 in 10000)	Patel, et al (2014)				
Insertive penile-vaginal intercourse	0.04% (4 in 10000)	Patel, et al (2014)				
Receptive oral intercourse	Low <sup>a</sup>	Varghese, et al. (2002); Page-Shafer, et al. (2002)				
Insertive oral intercourse	Low <sup>a</sup>	Varghese, et al. (2002)				
Other <sup>b</sup>						
Biting	Negligible	Deaths at al. (1999)				
Spitting	Negligible	Pretty, et al. (1999)				



# Blood and Body Fluid Pathogens Section: 6-40 – Human Immunodeficiency Virus (HIV) Infections Page 7 of 20 2018 10 01

Throwing body fluids (including semen or saliva)	Negligible
Sharing sex toys	Negligible

<sup>*a*</sup> HIV transmission through oral sex has been documented, but rare. Accurate estimates of risk are not available. It is prudent to recommend HIV post-exposure prophylaxis (PEP) for receptive oral sex with ejaculation, although discussion about the low risk should occur. Refer to Saskatchewan Guidelines for the Management of Blood and Body Fluids<sup>2</sup> for further consideration

<sup>b</sup> HIV transmission through these exposure routes is technically possible but extremely unlikely and cases are not well documented. Increased risk occurs when the activity involved exposure to blood

Source: New York State Department of Health AIDS Institute, 2013. AIDS (2014)

## **Risks for HIV Transmission**

- Multiple sexual partners (> 1 in 3 months).
- Unprotected sexual activity (i.e., no barrier protection).
- Sex with a person infected with HIV.
- Receptive anal/vaginal intercourse.
- Sharing of needles or other drug-using equipment.

## Specimen Collection and Transport

HIV infection is diagnosed by detection of antibodies, or of HIV antigens or nucleic acids in blood. For serological testing, collect blood in serum separator vacutainer (SST). Refer to Roy Romanow Provincial Laboratory (RRPL) Compendium of Tests at https://rrpltestviewer.ehealthsask.ca/. The serological test used at RRPL is the HIV combo assay, which detects the presence of both antibodies and the p24 antigen in serum. Reactive results in this assay are confirmed. See Saskatchewan HIV Testing Policy, Lab Testing Flowchart<sup>3</sup>.

# HIV viral load

Patients with confirmed HIV infection should have at least one HIV viral load assay performed. Refer to Roy Romanow Provincial Laboratory (RRPL) Compendium of Tests at https://rrpl-testviewer.ehealthsask.ca/.



<sup>&</sup>lt;sup>2</sup> <u>http://www.ehealthsask.ca/services/manuals/Pages/hiv-guidelines.aspx</u>

<sup>&</sup>lt;sup>3</sup> http://www.skhiv.ca/#!routine-testing/ciha

## HIV resistance genotyping

Patients who are receiving anti-retroviral therapy, and whose viral load increases should have a sample submitted for HIV resistance genotyping. Submit frozen specimens to RRPL with completed requisition for British Columbia Center of Excellence.

Newborns: sample referred to Reference laboratory for HIV detection by molecular methods.

# Public Health Investigation

I. Case

## <u>History</u>

Obtain as detailed a history as possible using the Attachment – HIV Data Collection Worksheet. This should be done in consultation and partnership with the ordering practitioner who initially diagnosed HIV in the individually. In order to monitor trends in epidemiology in Saskatchewan, it is important that all risk factors are asked and responses are documented. When a transmission risk is identified, timely follow-up must be completed.

- Inquire about factors that are associated with HIV <u>acquisition</u> or <u>transmission</u>:
  - Men who have sex with men (MSM);
  - multiple sexual partners;
  - injection drug use;
  - sharing injection or non-injection drug equipment; .
- history of sexual or needle-sharing contact with someone infected with HIV.
   Discuss all potential risks that the case has been exposed to with particular focus on parenteral exposures such as:
  - heterosexual sex with at risk individuals (person who injects drugs, men who have sex with men, persons from endemic country, injection drug use;
  - invasive body art (tattooing/piercing)<sup>4</sup>;
  - medical/dental procedures in sub-standard settings;
  - transfusions of blood/blood products in Canada;
  - transfusions of blood/blood products outside of Canada.

<sup>&</sup>lt;sup>4</sup> It is important to obtain details regarding dates of exposures and names/locations of the facilities in which exposures may have occurred. Consider whether investigation of any facility may be indicated. Consult with MHO. When personal service or medical/dental facilities are identified as a potential source for exposure, further investigation of other clientele may be warranted.



## **Public Health Interventions**

## Assessment

- It is important to know if the client is aware of their diagnosis or if the testing provider has not yet been able to notify the case. Prior to communicating with the client, discuss with health care provider who diagnosed the individual. Know whether the health care provider has informed their patient of the diagnosis and if they have collected information on contacts.
- Assess for contacts and obtain names and phone numbers of contacts as per Contact Investigation.

# Communication

- Individuals may be difficult to reach. Make several attempts to contact individuals using various methods (phone, text, home visit) at different times of the day. Some individuals' mobile service contracts only allow for text messaging. It is important to have policies and procedures that support the use of alternate modes of communication to assist in case follow-up.
- The primary care provider is an important partner in the public health follow-up of HIV. It is important to provide updates to care providers when referrals have been made to public health to assist in follow-up.

## Education

Providers are expected to be proficient in providing education in the topics below:

- Description of HIV infection progression, chronicity, treatment, management.
- Blood borne transmission/prevention, including risk reduction.
- *The Public Health Act, 1994*/Transmission/Prevention/Partner Notification of current and future partners:
  - the legal necessity to disclose HIV status with current and new sexual and needle-sharing partners.
- HIV post-exposure prophylaxis (PEP) use/availability in Saskatchewan.
- Contact notification responsibilities under *The Public Health Act, 1994*: Sexual/IDU/Other Blood Body Fluid Exposure.
- Infectious Diseases (ID) Specialist referral.

Education must be tailored to the individual and often requires repetition and reinforcement of learning. Information may need to be reinforced using written materials and repeated conversations.



## **Environmental Assessment**

• If personal service facilities are identified in the investigation, it may be prudent for a public health inspection to be made to ensure adequate infection prevention and control measures are in place.

# Exclusion

• Not applicable. Standard/Routine Infection Prevention and Control measures apply.

# Immunization

See Saskatchewan Immunization Manual – Chapter 7<sup>5</sup> for vaccines that HIV positive individuals are eligible for. Discuss with the regional medical health officer (MHO) and/or primary care practitioner/ID Specialist as required.

## Referrals

Cases should be referred to clinical and social services:

- Infectious Diseases (ID) specialist or other treating practitioner.
- Social programs as agreed to by client (e.g., community agencies that provide support to HCV positive people) or harm reduction programs for needle exchange services and related health services;
- Employee Health Department if case is a health care worker with a high risk of exposure to clients.
- Canadian Blood Services (CBS) if the case has a history of donation or receipt of blood or blood products. See <u>Appendix K – Notification to Canadian Blood</u> <u>Services.</u>
- Saskatchewan Transplant Program if the cases has a history of donation or receipt of tissues. See <u>Appendix M – Notification to the Saskatchewan</u> <u>Transplant Program.</u>
- In addition, a referral to an HIV Case Manager may be beneficial for clients that require additional supports.

# Testing

Cases should be advised that they should also be tested for other sexually transmitted and blood borne pathogens including chlamydia, gonococcal infections, syphilis, hepatitis B and hepatitis C.



<sup>&</sup>lt;sup>5</sup> http://www.ehealthsask.ca/services/manuals/Documents/sim-chapter7.pdf

# Treatment

The treatment of HIV infections is to be prescribed by an ID Specialist or General Practitioner mentored by an ID Specialist.

*Clinical management of cases involves follow-up testing which is not described in this document.* 

# II. Contacts/Contact Investigation Contact Definition

Contacts are defined as all sexual and needle/equipment-sharing partners of the case as well as others who may have been exposed to the case's blood or body fluids (e.g., trauma – see Mode of Transmission above) since:

- a. three months prior to the case's last negative HIV test **OR**
- b. onset of risk behaviour (for cases that have not been previously tested).
   In the case of "b", priority should be given to the most recent contacts.

All children born to mothers who are or may be HIV-infected need to be evaluated. Refer to Perinatal HIV Prevention Protocols<sup>6</sup>. This includes:

- a. children born within the window periods of the mother's positive test **AND**
- b. any children born since the last negative HIV test of the mother.

# Public Health Interventions

# Education

- Contacts should be identified and notified of their exposure to the disease.
- Contacts should be informed of their duties as outlined in the Disease Control Regulations:
  - to protect themselves by going to a physician or clinic nurse for testing and care;
  - to take all reasonable measures to reduce significantly the risk of infecting others.



<sup>&</sup>lt;sup>6</sup> https://skhiv.ca/pregnancy-and-newborn-care/

- Contacts should be assessed for risk behaviours and counselling should be provided to reduce risk exposures including the use of pre-exposure prophylaxis (PrEP).
- Referrals to harm reduction and supportive services should be provided as applicable.
- Contacts must be advised about blood and body fluid precautions while undergoing testing in the window period for HIV.

## Testing

- In addition to the education provided, pre-test counselling should be provided. *Canadian Guidelines on Sexually Transmitted Infections*<sup>7</sup> as well as the British Columbia Centre for Disease Control Communicable Disease Control Manual, Chapter 5: HIV Pre and Post Test Guidelines.
- The frequency and timing of testing should be based on the time since the most recent exposure/risk behaviour. Baseline testing is recommended at the time of contact notification. Follow-up tests should be conducted at 4 weeks and 3 months.
- If the exposure was 12 months ago, the baseline test would be all that is required unless the contact is engaging in other risk behaviours in which, case regular sexually transmitted and blood borne infection testing should be suggested.

# Prophylaxis

The Guidelines for Exposures to Blood and Body Fluids<sup>8</sup> outline the recommendations for the use of HIV post-exposure prophylaxis and the Saskatchewan Pre-Exposure Prophylaxis (PrEP) Guidelines<sup>9</sup> outline recommendations for PrEP. This may provide an opportune time to discuss PrEP for contacts that are engaged in ongoing exposures.

# Immunization

There is no vaccine to prevent HIV infections. Contacts should be provided immunizations as per the Saskatchewan Immunization Manual, Chapter 5<sup>10</sup> and Chapter 7.<sup>11</sup>



<sup>&</sup>lt;sup>7</sup> http://www.phac-aspc.gc.ca/std-mts/sti-its/cgsti-ldcits/section-5-8-eng.php.

<sup>&</sup>lt;sup>8</sup> http://www.ehealthsask.ca/services/manuals/Pages/hiv-guidelines.aspx

<sup>&</sup>lt;sup>9</sup> https://skhiv.ca/wp-content/uploads/2018/03/Pre-Exposure-Prophylaxis\_Guideline-Review-for-Primary-Care-Practitioners-in-Saskatchewan.pdf

<sup>&</sup>lt;sup>10</sup> http://www.ehealthsask.ca/services/manuals/Documents/sim-chapter5.pdf

<sup>&</sup>lt;sup>11</sup> http://www.ehealthsask.ca/services/manuals/Documents/sim-chapter7.pdf

## Exclusion

Not applicable. Standard blood and body fluid precautions apply until assured negative through testing as recommended above.

## III. Environment

## **Child Care Centre Control Measures**

Refer to the Saskatchewan Ministry of Health Infection Control Manual for Child Care Facilities.<sup>12</sup> All childcare centre staff should use standard precautions when handling all blood and body fluids. Children known to have HIV do not need to be excluded from childcare. If the child is known to bite, this should be discussed with the Medical Health Officer.

#### Institutional Control Measures

Refer to Saskatchewan Health Authority or former Regional Health Authority Infection Control Policies. Standard precautions should be followed by all staff working in health care settings. All health care settings should have policies and procedures in place for managing staff with occupational risk due to exposure to blood or body fluids. As well, there should be policies and procedures in place to manage occupational exposures to blood and body fluids.

For more information on occupational exposure see the Saskatchewan Guidelines for the Management of Exposures to Blood and Body Fluids.<sup>13</sup>

#### Personal Service Facilities

Refer to Saskatchewan Personal Service Facility Best Management Practices<sup>14</sup>.

 If personal service facilities are identified in the investigation, it may be prudent for a public health inspection to be made to ensure adequate infection prevention and control measures are in place. Consultation with the MHO is suggested.



<sup>&</sup>lt;sup>12</sup> http://www.saskatchewan.ca/live/births-deaths-marriages-and-divorces/starting-a-family/early-learning-and-child-care/child-care

<sup>&</sup>lt;sup>13</sup> http://www.ehealthsask.ca/services/manuals/Pages/hiv-guidelines.aspx

<sup>&</sup>lt;sup>14</sup> http://www.saskatchewan.ca/residents/environment-public-health-and-safety/environmental-health/personal-service-facilities

# Other Facilities with Alternate Caregivers and Other Residents (eg. group homes, foster homes, etc)

Standard precautions should be followed by all staff working in these settings. All settings should have policies and procedures in place for mitigating occupational risk of exposure to blood or body fluids. As well, there should be policies and procedures in place to manage occupational exposures to blood and body fluids should these occur.

For more information on occupational exposure see the Saskatchewan Guidelines for the Management of Exposures to Blood and Body Fluids.<sup>15</sup>

## **IV. Epidemic Measures**

When two or more cases occur in association with a common exposure, search for additional cases. Screen contacts and implement measures to interrupt further transmission as appropriate to the situation

Medical Health Officers may declare and outbreaks of HIV that has been identified through contact tracing efforts. Responding to an HIV or HCV outbreak may require augmenting and redirecting resources, engaging a large and diverse group of partners and stakeholders, building upon collaborations and developing targeted communication messages for specific groups. Increased resources are usually needed to respond to the increased number of new diagnoses and to identify the root causes of the outbreak. Refer to the US CDC publication, *Managing HIV and Hepatitis C Outbreaks Among People Who Inject Drugs*<sup>16</sup> for reference.

# **Prevention and Education**

Refer to the Blood and Body Fluid Pathogens Introduction and General Considerations section of the manual that highlights topics for client education that should be considered.

<sup>&</sup>lt;sup>15</sup> http://www.ehealthsask.ca/services/manuals/Pages/hiv-guidelines.aspx

 $<sup>^{16}\</sup> https://www.cdc.gov/hiv/pdf/program resources/guidance/cluster-outbreak/cdc-hiv-hcv-pwid-guide.pdf$ 

Health education efforts should include both broad-based campaigns to raise awareness of risk, modes of transmission, and prevention measures, and reduce stigma as well as targeted programs to educate and reduce risk in target populations.

Routine testing should be promoted by health care providers. Refer to the Public Health Agency of Canada HIV Screening and Testing Guide<sup>17</sup> and the SK HIV Testing Policy<sup>18</sup> for additional information on routine testing.

## Immunization

There is no immunization available for the prevention of HIV infection.

## Pre-Exposure Prophylaxis

PrEP is an important prevention intervention that should be offered as part of an overall risk reduction strategy. PrEP involves the use of antiretroviral medications by confirmed HIV negative individuals with ongoing risk of HIV acquisition. It is initiated before HIV exposures. It should be used in conjunction with behavioural risk counselling and other harm reduction interventions. Refer to the Saskatchewan Pre-Exposure Prophylaxis Guidelines.<sup>19</sup>

## Education

- Health education efforts should include both broad-based campaigns to raise awareness of risk, modes of transmission, and prevention measures, and reduce stigma as well as targeted programs to educate and reduce risk in at-risk populations.
- Personal service providers should be referred to Saskatchewan Personal Service Facility Best Management Practices<sup>8</sup> for infection prevention and control measures.

<sup>18</sup> http://www.skhiv.ca/#!routine-testing/ciha



<sup>&</sup>lt;sup>17</sup> http://www.phac-aspc.gc.ca/aids-sida/guide/hivstg-vihgdd-eng.php

<sup>&</sup>lt;sup>19</sup> https://skhiv.ca/wp-content/uploads/2018/03/Pre-Exposure-Prophylaxis\_Guideline-Review-for-Primary-Care-Practitioners-in-Saskatchewan.pdf

## Revisions

Date	Change
September 2018	<ul> <li>Clarified the purpose for notification of cases to public health</li> <li>Incorporated Stages of HIV based on CD4 counts</li> <li>Incorporated a placeholder for an Epidemiology and Occurrence section to the chapter.</li> <li>Removed case definition for AIDS as it is included in its own chapter.</li> <li>Incorporated standardized HIV Data Collection Worksheet.</li> <li>Rearranged and updated the style into the new format of the Manual.</li> <li>Added information on U=U and PrEP.</li> <li>References reviewed and updated as applicable.</li> </ul>



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#### **HIV Notification Form**

Please complete all sections



Panorama QA complete: Yes No Initials:

#### A) PERSON REPORTING - HEALTH CARE PROVIDER INFORMATION

Clinic Name:	FOR PUBLIC HEALTH OFFICE USE ONLY:
Location:	Service Area:
Attending Physician or Nurse:	Date Received:
Address:	Panorama Client ID:
Phone number:	Panorama Investigation ID:

#### B) CLIENT INFORMATION

Last Name:	First Name: and Middle Name:	Alternate Name:
DOB: YYYY / MM / DD Age: Health Card Province: Health Card Number (PHN):	Gender: Male Female Unknown Other <u>Gender Identity:</u> Transgender Male-to-female Transgender Female-to-male Undifferentiated Other (specify)	Phone : Primary Home: Mobile contact: Workplace: Alt Contact: Name: Relationship:
Place of Employment/School:	Email Address:	Preferred Communication Method:
Address Type:	□ Primary Home □ Temporary	□ Legal Land Description
C) IMMIGRATION INFORMATION		
Country Born In:		

#### D) DISFASE EVENT HISTORY

Country Emigrated from: \_\_\_\_

Site / Presentation: Adults, adolescents, and children > 18 months		□ Children <18	months		
Staging (see CDC Man	ual): 🛛 Stage 0	□ Stage 1 (CD4 ≥500)	□ Stage 2 (CD4 200-499)	□ Stage 3 (CD4 <200)	🗆 Unknown

Arrival Date: YYYY / MM / DD

OR Arrival Year YYYY

E) SIGNS & SYMPTOMS

	YES	NO		YES	NO	SPECIFY
Asymptomatic			Symptoms prior to or at time of testing?			
Initial CD4 result						

#### **HIV Notification Form**

Please complete **all** sections

#### F) RISK FACTORS (Please complete <u>all</u> Risk Factors from 3 months prior to last known negative result –specify dates as needed) Legend: N-No, NA-Not Asked, U-Unknown

DESCRIPTION	Yes Start date	N, NA, U	Add'l Info		
Sexual Behaviour – MSM +	TE	.,-			
Sexual Behaviour - Heterosexual Sex	TE				
Sexual Behaviour - Heterosexual sex with person who injects drugs	TE				
Sexual Behaviour - Heterosexual sex with MSM	TE				
Sexual Behaviour - Heterosexual sex with person with hemophilia/coagulation disorder	TE				
<b>Sexual Behaviour</b> - Heterosexual sex with person from endemic country (Add'I Info)					
Sexual Behaviour – Heterosexual sex with person with confirmed/suspected HIV/AIDS (Add'I Info)	YYYY / MM/DD				
Sexual Behaviour – Sex with a known case	YYYY/MM/DD				
Sexual Behaviour - Unknown/Anonymous Partner (Add'l Info)	TE				
Sexual Behaviour - E-partnering internet/apps (Add'l Info.)	TE				
Sexual Behaviour - Goods provided (food,	TE				
shelter, money or drugs) in exchange for sex Sexual Behaviour - Goods received (food, shelter,	TE				
money or drugs) in exchange for sex					
Sexual Behaviour - Events with multiple sexual partners (Add'I Info)	TE				
<b>Exposure</b> - Blood and body fluids (not otherwise listed) (Add'l Info.)	YYYY / MM/DD				
<b>Exposure</b> - Invasive body art (e.g. tattoo, body piercing, scarification)	YYYY / MM/DD				
<b>Exposure</b> - Non medical, non-occupational source	YYYY/MM/DD				
(acupuncture, breastmilk) (Add'l Info) <b>Exposure</b> - Occupational - HIV contaminated	YYYY / MM/DD				
blood, body fluid					
Special Population - Infant born to an infected	YYYY/MM/DD				
mother		ļ			
Special Population - From or residence in an endemic country (Add'I Info)					
Special Population – Pregnancy					
Special Population - Self-reported Indigenous					
Substance Use - Injection drug use (including steroids)	YYYY / MM/DD				
Risk Behavior - Sharing injection drug equipment	YYYY / MM/DD TE				
Medical Treatment - Blood, blood product or tissue recipient (Add'l Info.)	YYYY / MM/DD INTERVENTION				
Medical Treatment - Other (transplant, surgery, dental, oscopy, etc.) (Add'l Info)	YYYY / MM/DD INTERVENTION				
Blood, blood product, tissue or transplant <b>donor</b>	or Document referral in Interventions and complete Appendix K – Referral to CBS, and upload into Document Management				
Unable to obtain Risk Factors $\Box$ yes (not entered in Panorama – update in disposition)					

#### G) UNKNOWN/ANONYMOUS CONTACTS

Anonymous contacts: \_\_\_\_\_ (number of contacts that the individual cannot name)

#### Include known contacts on the following pages





#### **HIV Notification Form - Contacts**

Please complete all sections.

Case Name:

Please include information on additional contacts on a separate sheet

Page \_\_\_\_\_ of \_\_\_\_\_

CONTACTS						
Last Name: Fir	rst Name: and Middle Name	::	Alternate Nam	ie:		
DOB: YYYY / MMM / DD Age: Ge	ender: 🗆 Male 🗖 Female	🗆 🗆 Unknown	□ Other			
Phone #:  Primary Home:		e-mail Address	:			
□ Workplace: □ Mobile contact:						
alternate phone: Relationship:						
Online Names:						
Site/Service:	User Name:					
Place of Employment/School:		Is contact preg	nant?	🗆 Yes 🗖 No	Unknown	
		Is contact HIV	ositive	🗆 Yes 🗖 No	🗆 Unknown	
		If yes, did they	inform case?	🗆 Yes 🗖 No	🗆 Unknown	
Address Type: 🗆 No fixed 🗆 Postal Address 🗆 P	Primary Home Tem	porary 🗖 Legal I	Land Description			
Mailing (Postal address):						
Street Address or FN Community (Primary Home):	Street Address or FN Community (Primary Home):					
Exposure Dates: 1st YYYY / MMM / DD to YYY	YY / MMM / DD					
Exposure Type:  Heterosexual  Sharing Injection Drug E	Equipment 🗖 MSM					
Comments:	INTERVENT	ION				
	Testing	□ Advised □	Received 🛛 🖬	Referral (Specify)		

#### CONTACTS

Last Name:	First Name: and Middle Nam	e:	Alternate Name		
DOB: YYYY / MMM / DD Age:	Gender: 🗆 Male 🛛 Femal	e 🗖 Unknown	□ Other		
Phone #:  Primary Home:  Workplace:  Mobile contact:  alternate phone:  Relationship:		e-mail Address	:		
Online Names:					
Site/Service:	User Name:				
Place of Employment/School:		Is contact preg	nant?	🗆 Yes 🗖 No	🗆 Unknown
		Is contact HIV	oositive	🗆 Yes 🗖 No	🗆 Unknown
		If yes, did they	inform case?	🗆 Yes 🗖 No	🗆 Unknown
Address Type:  No fixed  Postal Address	□ Primary Home □ Ten	porary 🗆 Legal	Land Descriptior	1	
Mailing (Postal address):					
Street Address or FN Community (Primary Home):					
Exposure Dates: 1st YYYY / MMM / DD to	YYYY / MMM / DD				
Exposure Type:  Heterosexual  Sharing Injection Dr	rug Equipment 🛛 MSM				
Comments:	INTERVEN	ΓΙΟΝ			
	Testing	□ Advised □	Received 🛛	Referral (Specify)	



#### HIV – Public Health Follow-Up

Panorama QA complete: 
QYes 
No Initials:



Panorama Client ID: Panorama Investigation ID: \_

YYYY / MM / DD

#### A) CLIENT INFORMATION

A) CLIENT INFORMATION	LHN ->SUBJE	CT -> CLIENT DETAILS -> PERSONAL INFORMATION
Last Name:	First Name: and Middle Name:	Alternate Name:
DOB: YYYY / MM / DD Age:	Gender: □ Male □ Female □ Unknown □ Other	PHN:

Disease Summary Classification: CASE:	Date	Classification: CONTACT:	Date	LAB TEST INFORMATION:
Lab Confirmed	yyyy / MM / DD	🗖 Contact	YYYY / MM / DD	Date specimen collected:
□ Suspect	yyyy / MM / DD	□ Not a Contact	YYYY / MM / DD	YYYY / MM / DD
Person Under Investigation	yyyy / MM / DD	$\Box$ Person Under Investigation	YYYY / MM / DD	
Disposition: FOLLOW UP:		<u></u>		
In progress	yyyy / MM / de	Complete		yyyy / MM / DD
Incomplete - Declined	yyyy / MM / de	Not required		yyyy / MM / DD
Incomplete – Lost contact	yyyy / MM / de	🗆 🗖 Referred – Out o	of province	yyyy / MM / DD
□ Incomplete – Unable to locate	yyyy / MM / de	(Specify where)		YYYY / MM / DD

Intervention Type and Sub Type:			VILIVLIVIION3-2INTERVEI	1101/ 201
Assessment:		Immunization: Investigator name		
□ Assessed for contacts Investigator name	YYYY / MM / DD	Eligible Immunization recommended	yyyy / MM	/ DD
□ Client aware of diagnosis Investigator name	YYYY / MM / DD	Immunization nurse notified	yyyy / MM	/ DD
Communication:	, ,	Environmental health:		
Phone call (morning)     Investigator name	YYYY/ MM/ DD	Personal Service Facility inspection	yyyy / mm	/ DD
□ Phone call (afternoon) Investigator name	YYYY/ MM/ DD	Investigator name		
□ Phone call (evening) Investigator name	YYYY/ MM/ DD	Referral: Investigator name		
□ Text Message sent Investigator name	YYYY/ MM/ DD	Canadian Blood Services	yyyy / MM	
E-mail     Investigator name	YYYY/ MM/ DD	□ Child Protective Services	YYYY / MM	
□ Home visit Investigator name	YYYY/ MM/ DD	Harm Reduction Services	yyyy / MM	/ DD
□ Letter Sent Investigator name	YYYY/ MM/ DD	HIV Case Management	yyyy / MM	,
□ Letter (See Document Management)	YYYY/ MM/ DD	□ Infectious Disease Specialist	YYYY / MM	
Investigator name		<ul> <li>Primary Care Provider</li> <li>Saskatchewan Transplant Program</li> </ul>	yyyy / mm yyyy / mm	
Ordering practitioner contacted	YYYY/ MM/ DD	$\Box$ Consultation with MHO	YYYY / MM	,
Investigator name			,	/
Other communication (See Investigator Notes)	YYYY/ MM/ DD	Other:		
Investigator name				1.00
5		Other (specify)	YYYY / MM	/ DD
General: Investigator name		Other Investigation Findings	YYYY/ MM ,	/חח
Disease-Info/Prev-Control	YYYY/ MM / DD	See Document Management	YYYY/ MM	
Disease-Info/Prev-Cont/Assess'd for Contacts	YYYY/ MM / DD		,	
Education/counselling:		Testing:		
□ Prevention/Control measures Investigator name	e yyyy / MM / DD	Laboratory testing recommended	yyyy / MM	
Disease information provided Investigator name	e yyyy / MM / DD	□ STBBI Testing recommended -See Inves	stigator Notes YYYY / MM	/ DD
Other (See Investigator Notes)	YYYY / MM / DD			
Date Intervention subtype Comm	nents		Next follow-up Date	Initials
YYYY / MM / DD			YYYY / MM / DD	
YYYY / MM / DD YYYY / MM / DD			YYYY / MM / DD YYYY / MM / DD	

YYYY / MM / DD

#### **HIV Public Health Follow-up**

Please complete all sections.

#### **D)** OUTCOMES (Optional except for severe influenza)

LHN-> INVESTIGATION-> OUTCOMES

Other     Fatal YYYY/N	YYYY/MM/DD /M/DD	Cause of Death: (if Fatal wa	s selected)		
) Transmission Even	ts	LHN -> INVESTIGATION	I-> EXPOSURE SUMMARY -> TRAN	ISMISSION EVENT SUMMA	ARY -> QUICK E
Transmission	Exposure Name	Setting type		Date/Time	# of
Event ID		Important:		(included the earliest	contacts
(system-generated can		· · · · · ·	e setting for the TE; if >1 select	transmission date to the latest date)	
be documented below)	UN/ Contract Jaw ID #	multiple settings)			
	HIV Contact – Inv ID #	□ Sexual Exposure			
		Type of community conta	ct (IDU)		
		Public facilities	Multiple Settings		

#### F) Total number of contacts

LHN -> INVESTIGATION-> EXPOSURE SUMMARY -> TRANSMISSION EVENT SUMMARY -> TE HYPERLINK -UNKNOWN/ANONYMOUS CONTACTS

(Total number of unknown and known contacts)

nitial Report	Date initial report completed:
completed by:	YYYY / MMM / DD

#### CONTACTS

Last Name:	First Name: and Middle Name:		Alternate Name:
DOB:         YYYY         MMM / DD         Age:           HSN:	Gender: 🗆 Male 🛛 Female	Unknown	□ Other
Phone #:  Primary Home: Workplace: Mobile contact: alternate phone: Relationship:		e-mail Address:	
Online Names:			
Site/Service:	User Name:		
Place of Employment/School:		Is contact pregna	ant? 🛛 Yes 🗆 No 🗖 Unknown
		Is contact HIV po	sitive 🗆 Yes 🗆 No 🗖 Unknown
		If yes, did they in	iform case? 🛛 Yes 🗆 No 🗖 Unknown
Address Type:  No fixed  Postal Address	□ Primary Home □ Temp	orary 🛛 Legal La	nd Description
Mailing (Postal address): Street Address or FN Community (Primary Home):			
Exposure Dates: 1st YYYY / MMM / DD to	YYYY / MMM / DD		
Exposure Type:  Heterosexual  Sharing Injection D	rug Equipment 🛛 MSM		
Comments:	INTERVENTIO	ON	
	Testing 🗆	□ Advised  □ R	eceived 🛛 🗖 Referral (Specify)

#### Complete more contact sheets if needed

Please see the following pages for the AIDS Case Report Form.



Public Health Agence de santé				Protected when completed
Agency of Canada publique du Canada	For p	provincial/territorial use		For use by PHAC
HIV/AIDS Case Report Adult, Adolescent and Pediatric		ncial/territorial ID Number		EPIC No.
(non maternal-fetal) Cases	D	en e		
HIV AIDS New case report Update	Provir	nce/Territory to which case is attributed		Date received YY MM DD
Reporting physician's name		City	Telephone	number
			( )	
Hospital or clinic		City	Province/Te	erritory
Is another physician providing ongoing care to this patient? Yes	No	If so, please provide name, city and telephone City	number. Telephone r	number
			( )	
Patient's initials     Sex     Date of birth       First     Middle     Last       M     F	Vital Statu	IS Alive (If yes, date last known to be alive) Dead (If yes, date of death)	YY M	
• Is the patient: (please ask patient to assist you in answering this of	questic	on)		
<ul> <li>White</li> <li>Black (e.g. African, Haitian, Jamaican, Somali, etc.)</li> <li>North American Indian Métis Inuit</li> <li>Asian (e.g. Chinese, Japanese, Vietnamese, Cambodian, Indonesian, Laotian, Korean, Filipino, etc.)</li> </ul>		South Asian (e.g. East Indian, Pakistani, Sri Arab/West Asian (e.g. Armenian, Egyptian, Latin-American (e.g. Mexican, Central/Souti Other – includes mixed ethnicity (specify	Iranian, Lebai h American, e	nese, Moroccan, etc.)
What language does this person speak most often at home? C	ountry	of birth		Year of arrival in Canada
	Can	ada 📃 Other (specify) 🔶		
City and province/territory of residence at diagnosis		Current city and province/territory of	rosidonco	
City and province/territory of residence at diagnosis           City         Province/Territory           First 3 digits of Post	stal Cod	Current city and province/territory of r City Province/Territory	esidence	First 3 digits of Postal Code
		_		
SECTION II – RISK(S) ASSOCIATED WITH THE TRANSMISS	SION	OF HIV IN THIS PATIENT		
Since January 1978 and preceding the diagnosis of HIV/AIDS, thi Yes No Unknown	is patie	ent had: (check ALL that apply)		
Sex with a male.				
Sex with a female.				
Heterosexual sex with: (check ALL that app	ly)			
• an injection drug user;				
• a bisexual male;				
a transfusion recipient with documente	ed HIV i	nfection;		
a person with hemophilia/coagulation of the second se	disorde	r,		
		al transmission predominates. If yes, specify co	-	
		ection or AIDS (whether or not risk factor is know	wn).	
	or IX for	r treatment of hemophilia/coagulation disorder.		
	d comp	onents such as packed red cells, plasma, platel	lets or cryopre	cipitate.
L If yes, please complete Section 2 of the S Exposure to HIV-contaminated blood or body	y fluids	or concentrated		
Virus in an occupational setting. <b>If yes, spec</b>	-	•		
If yes, please give details in Section VI "A				
		Id have been the source of the infection (e.g. ac ate and location in Section VI "Additional Inf		
Since January 1978, has this patient donated blood, plasma, platelets, organ <b>If yes, please give details of type of donation, date and location in Sect</b> Has the Red Cross or other appropriate donor program been notified? Do you want a public health official to ensure this notification?			Yes	No Unknown
PHAC/ASPC 4205 F (03-2006) Distribution: White – Medical Officer of Health Yellow – Ministry	/ of Hea	lth Pink – PHAC		

SECTION III - LABORATORY DA	ATA							
Does this case have evidence, as a	lefined in the abo	ve instructions, of	Date of	first positive	e HIV test (if	known)	Current CD4 count (if ki	nown)
HIV infection? Yes No Unknow	wn			Year	Month			cells/µ I
SECTION IV – DISEASES INDIC	ATIVE OF AIDS				1			
DISEASES	-	Diagnostic method		DISEA	SES		Date of Diagnosis Diagnos	
Bacterial pneumonia, recurrent	Year Month	Definitive Presumptive		vcohacterium	avium compl	ev or	Year Month Definitive	e Presumptive
Candidiasis (bronchi, trachea or			M.	kansasii	or extrapulmo			
lungs)			My	cobacterium	of other spec			
Candidiasis (esophageal)				identified spe				
Cervical cancer, invasive			(di	tuberculosis sseminated c	or extrapulmo	nary)		
Coccidioidomycosis (disseminated or extrapulmonary)			(P	Specify Si	ete SECTION	i V)		
Cryptococcosis (extrapulmonary)				Miliary		Pleurisy	y Other respi	ratory
Cryptosporidiosis (chronic intestinal, >1 mo. duration)				C.N.S.	. [	Bone a	nd joint 🗌 Genitourina	ary
Cytomegalovirus disease (other than in liver, spleen or nodes)				Other	(specify) →	•		
Cytomegalovirus retinitis (with loss of vision)			<i>М.</i> (Р	tuberculosis	(pulmonary) ete SECTION	I V)		
Encephalopathy, HIV-related (dementia)			Pr	eumocystis c	<i>arinii</i> pneumo	onia		
Herpes simplex: chronic ulcer(s)				ogressive mu ikoencephalo				
(>1 mo. duration) or bronchitis, pneumonitis or esophagitis					ticemia, recu	rrent		
Histoplasmosis (disseminated or extrapulmonary)				xoplasmosis				
Isosporiasis, chronic intestinal (>1 mo. duration)			Wa	asting syndro	me due to HI	V		
Kaposi's sarcoma			Di	seases affe	ecting pedi	atric cas	es only (<15 years old)	)
Lymphoma, Burkitt's (or equivalent term)					ons, multiple Iding recurrer			
Lymphoma, immunoblastic			pn	eumonia)	U U			
(or equivalent term) Lymphoma, primary in brain					stitial pneumo phoid hyperpl			
SECTION V – TUBERCULOSIS								
1. Before the diagnosis of AIDS, was tuberculosis?	s this patient ever	treated for	Y	es – when? -	→ Year	Mon	th No [	Unknown
2. Has this patient ever had a PPD s	kin test?	Yes	- What wa	as the size in r	mm? → [		mm No	Unknown
3. If the PPD test was negative, was		y tested? Yes	No	Unknowr	n If yes, were	any sites pos	sitive? Yes No [	Unknown
SECTION VI – ADDITIONAL INFO	ORMATION OR	COMMENTS						
(Please use this section for info			isition	of the virus	s, etc.)			
Person completing this form				Teleph	one number		Date report comp	
				(	)			
FOR PROVINCIAL/TERRITORIAL US	E: To which expo	sure category has th	is patient	been assig	ned?			
Men who have sex with men (MSM)			SM and IDU		_	al – Endemic		exual
Blood transfusion recipient	Clotting fact	tor recipient Oc	cupational	exposure	Heterosexu	al – Partner a	at risk NIR – Other	

PHAC/ASPC	4205	F	(03-2006)	



Panorama QA complete: □Yes □No Initials:

#### **AIDS Data Collection Worksheet**

Please complete all sections.

PANORAM

Panorama Client ID: \_ Panorama Investigation ID: \_

A) CLIENT INFORMATION	
-----------------------	--

LHN ->SUBJECT -> CLIENT DETAILS -> PERSONAL INFORMATION

A CLIENT INFORMATION		LHIN	->20DJEC1 ->	CLIENT DETAILS -> PERSONAL INFORIVIATION
Last Name:	First Name: and N	/liddle Name:	Alternate N	lame:
DOB: YYYY / MM / DD Ag Health Card Province: Health Card Number (PHN): Place of Employment/School:	ge: <u>Gender: _</u>	emale-to-male	Phone : Primary Home: Mobile contact: Workplace: Alt Contact: Name: Relationship:	
Place of Employment/School:	Email Address:			ommunication Method: ] Work 🗖 E-mail 🗖 Text
Address Type: No fixed Postal Address Mailing (Postal address): Street Address or FN Community (Prim Address at time of investigation if not	nary Home):	Temporary □Lega	al Land Descrip	ption
<b>B)</b> INVESTIGATION INFORMATION		SUBJECT SUMM	ARY->STBBI EI	NCOUNTER GROUP->CREATE INVESTIGATION
Disease Summary Classification: CASE:	Date	Investigation Information Disposition:	n	Date
Confirmed	YYYY / MMM / DD	Complete	vince	yyyy / mmm / dd
FOLLOW UP:         In progress         Incomplete - Declined         Incomplete - Lost contact         Incomplete - Unable to locate	YYYY / MM / DD YYYY / MM / DD YYYY / MM / DD YYYY / MM / DD	<ul> <li>Complete</li> <li>Not required</li> <li>Referred – Out of pro (Specify where)</li> </ul>	vince	YYYY / MM / DD YYYY / MM / DD YYYY / MM / DD YYYY / MM / DD
<b>REPORTING NOTIFICATION:</b> Name of Attending Physician or Nurse	:	Location:		
Provider's Phone number:		Date Received (Public He	ealth): YYYY	/ MMM / DD

□ Other\_ Type of Reporting Source: 🗖 Health Care Facility □ Nurse Practitioner □ Physician C) OUTCOMES (optional except for severe influenza, LHN-> INVESTIGATION-> OUTCOMES □ Not yet recovered/recovering YYYY / MM / DD □ ICU/intensive medical care \_ YYYY / MM / DD □ Hospitalization YYYY / MM / DD □ Recovered □ Intubation /ventilation yyyy / MM / DD 🗆 Unknown YYYY / MM / DD YYYY / MM / DD 🗆 Fatal YYYY / MM / DD Other YYYY / MM / DD

Cause of Death: (if Fatal was selected)

# **AIDS Data Collection Worksheet**

Please complete all sections

Panorama Client ID: \_\_\_\_\_ Panorama Investigation ID: \_\_\_\_\_

#### DISEASES INDICATIVE OF AIDS

DESCRIPTION	Date of Diagnosis	Definitive	Presumptive
Bacterial pneumonia, recurrent	,,		
Candidiasis (bronchi, trachea or lungs)			
Candidiasis (esophageal)			
Cervical cancer, invasive			
Coccidiodomycosis (disseminated or extrapulmonary)			
Cryptococcosis (extrapulmonary)			
Cryptococcosis (chronic intestinal, >1mo. Duration)			
Cytomegalovirus disease (other than in liver, spleen or nodes)			
Cytomegalovirus retinitis (with loss of vision)			
Encephalopathy, HIV-related (dementia)			
Herpes simplex: chronic ulcer(s) (>1 mo. Duration)or bronchitis, pneumonitis or esophagitis			
Histoplasmosis (disseminated or extrapulmonary)			
Isoporiasis, chronic intestinal (>1mo. Duration)			
Kaposis's sarcoma			
Lymphoma, Burkitt's (or equivalent term)			
Lymphoma, immunoblastic (or equivalent term)			
Lymphoma, primary in brain			
Mycobacterium avium complex or M. kansasii (disseminated or extrapulmonary)			
Mycobacterium of other species or unidentified species			
<i>M. tuberculosis</i> (disseminated or extrapulmonary) <b>Specify in comments</b> : <b>Millary, Pleurisy, Other respiratory, CNS, Bone and Joint, Genitourinary</b> )			
M. tuberculosis (pulmonary)			
Pneumocystis carinii pneumonia			
Progressive multifocal leukoencephalopathy			
Salmonella septicemia, recurrent			
Toxoplasmosis of brain			
Wasting syndrome due to HIV			
<15 years of age – Bacterial infections, multiple or recurrent (excluding recurrent bacterial pneumonia)			
<15 years of age – Lymphoid interstitial pneumonia and/or Pulmonary lymphoid hyperplasia			

#### Additional Information or Comments: