

MHSU 6781 - PROVIDER REPORT FORM FOR SEXUALLY TRANSMITTED AND BLOOD-BORNE INFECTIONS (STBBI) AND STI TREATMENT



NEW REPORT _____ (YYYY-MM-DD) UPDATED REPORT _____ (YYYY-MM-DD)

I. CLIENT IDENTIFICATION

subject > client details > personal information

| | | | | | | |
|--|--|---|--|------------------------------------|-------------------------------------|--|
| LAST NAME | | FIRST NAME | | DATE OF BIRTH (YYYY-MM-DD) | | |
| SEX <input type="checkbox"/> FEMALE <input type="checkbox"/> INTERSEX <input type="checkbox"/> MALE <input type="checkbox"/> UNKNOWN | | GENDER IDENTITY (VOLUNTARY, SELF-REPORTED) <input type="checkbox"/> CISGENDER (SAME AS SEX AT BIRTH) <input type="checkbox"/> TRANSGENDER PERSON <input type="checkbox"/> DECLINED <input type="checkbox"/> TRANSGENDER MAN <input type="checkbox"/> TRANSGENDER WOMAN <input type="checkbox"/> OTHER (SPECIFY) | | | AGE (YRS) (IF DOB NOT COMPLETED) | |
| REGISTRATION NUMBER (FORMER MHSC) 6 DIGITS | | HEALTH NUMBER (PHIN) 9 DIGITS | | ALTERNATE ID SPECIFY TYPE OF ID | | |
| ADDRESS AT TIME OF DIAGNOSIS → <input type="checkbox"/> ADDRESS IN FIRST NATION COMMUNITY | | | | | CITY/TOWN/VILLAGE | |
| PROVINCE/TERRITORY | | POSTAL CODE (A#A #A#) | | PHONE NUMBER (### - ### - ####) | | |
| ALTERNATE IDENTIFYING OR LOCATION INFORMATION (IF ANY. E.G. ALTERNATE NAME, SOCIAL MEDIA, ALTERNATE ADDRESS) | | | | | | |
| PREVIOUS NON-NOMINAL CODE(S) OR NAME(S) USED FOR POSITIVE HIV TESTS IF APPLICABLE (SPECIFY COUNTRY/PROVINCE, CODE/NAME, AND DATES YYYY-MM-DD IF KNOWN) | | | | | | |

II. PREGNANCY

subject > risk factors

IS CLIENT PREGNANT/POST PARTUM? YES EDD OR DELIVERY DATE: YYYY-MM-DD NO UNKNOWN

III. INFECTION INFORMATION

investigation > investigation details > disease summary > update > disease event history

| | | | | | | | |
|--|--|--|------------------------------|---|--------------------------------------|------------------------------|-----------------------------------|
| REASON FOR REPORTING: | | <input type="checkbox"/> LAB CONFIRMED INFECTION(S) (SPECIFY BELOW) | | <input type="checkbox"/> STBBI TREATMENT PROVIDED (CONTACTS OR CLINICAL CASES) (TEST RESULTS PENDING OR NOT DONE) PROCEED TO TREATMENT INFORMATION | | | |
| LAB CONFIRMED INFECTIONS (CHECK ALL THAT APPLY) | <input type="checkbox"/> CHLAMYDIA <input type="checkbox"/> GONORRHEA | <input type="checkbox"/> CHANCROID | <input type="checkbox"/> LGV | <input type="checkbox"/> HEPATITIS B | <input type="checkbox"/> HEPATITIS C | <input type="checkbox"/> HIV | <input type="checkbox"/> SYPHILIS |
| SPECIMEN COLLECTION DATE (YYYY-MM-DD) | | | | | | | |

IV. TREATMENT INFORMATION

investigation > prescriptions > prescription summary

| | | | | | | | |
|--|---|--|--|--|--|--------------------------|--|
| PRESCRIBER NAME | | PRESCRIBER/TREATMENT FACILITY | | | | | |
| SYPHILIS | <input type="checkbox"/> BENZATHINE PENICILLIN G 2.4 million units, IM, 1 dose START DATE (YYYY-MM-DD): | <input type="checkbox"/> BENZATHINE PENICILLIN G 2.4 million units, IM weekly, 2 doses START DATE (YYYY-MM-DD): | <input type="checkbox"/> BENZATHINE PENICILLIN G 2.4 million units, IM weekly, 3 doses START DATE (YYYY-MM-DD): | <input type="checkbox"/> CEFTRIAXONE 1 g daily x 10 days, IV / IM (circle one) START DATE (YYYY-MM-DD): | | | |
| | <input type="checkbox"/> CEFTRIAXONE 2 g daily x 10 days, IV / IM (circle one) START DATE (YYYY-MM-DD): | <input type="checkbox"/> DOXYCYCLINE 100 mg PO BID x 14 days START DATE (YYYY-MM-DD): | <input type="checkbox"/> DOXYCYCLINE 100 mg PO BID x 28 days START DATE (YYYY-MM-DD): | <input type="checkbox"/> PENICILLIN G 3 - 4 M IV Q4H x 10-14 days START DATE (YYYY-MM-DD): | | | |
| CHLAMYDIA, GONORRHEA | <input type="checkbox"/> AZITHROMYCIN 1g PO, single dose START DATE (YYYY-MM-DD): | <input type="checkbox"/> CEFIXIME 800 mg PO, single dose START DATE (YYYY-MM-DD): | <input type="checkbox"/> DOXYCYCLINE 100 mg PO BID x 7 DAYS START DATE (YYYY-MM-DD): | <input type="checkbox"/> METRONIDAZOLE 500 mg PO BID x 14 DAYS START DATE (YYYY-MM-DD): | | | |
| | <input type="checkbox"/> AMOXICILLIN 500 mg PO TID x 7 DAYS START DATE (YYYY-MM-DD): | <input type="checkbox"/> CEFTRIAXONE 250 mg IM, single dose START DATE (YYYY-MM-DD): | <input type="checkbox"/> ERYTHROMYCIN 500 mg PO QID x 7 DAYS START DATE (YYYY-MM-DD): | | | | |
| OTHER TREATMENT (LGV OTHER INFECTION, IF APPLICABLE) | SPECIFY: | | | | | START DATE (YYYY-MM-DD): | |
| UPDATE TO PREVIOUS INFORMATION SUBMITTED | <input type="checkbox"/> SPECIFY DETAILS ON ANY CHANGE TO SYPHILIS TREATMENT PLAN (E.G. CLIENT DID NOT ATTEND FOR ANOTHER DOSE) | | | | | | |
| | <input type="checkbox"/> PREVIOUSLY REPORTED TREATMENT PRIOR TO POSITIVE TEST – PROVIDE FORM COMPLETION DATE (YYYY-MM-DD) | | | | | | |

CONFIDENTIAL – WHEN COMPLETED

| | |
|-------------------|-----------------------------------|
| CLIENT LAST NAME: | CLIENT FIRST NAME: |
| PHIN: _____ | : _____ OR _____ DOB (YYYY-MM-DD) |

V. PRESENTATION/STAGING (FOR LAB CONFIRMED CASES ONLY)

[investigation](#) > [investigation details](#) > [investigation information](#)

| | | | |
|--|--|--|---|
| COMPLETE FOR CHLAMYDIA, GONORRHEA, LGV, CHANCROID ONLY | | | |
| PRESENTATION | | | |
| <input type="checkbox"/> ARTHRITIS | <input type="checkbox"/> GENITAL | <input type="checkbox"/> PHARYNGEAL | <input type="checkbox"/> RECTAL/ANAL |
| <input type="checkbox"/> EYE | <input type="checkbox"/> PELVIC INFLAMMATORY DISEASE | <input type="checkbox"/> PNEUMONIA | <input type="checkbox"/> OTHER (SPECIFY): |
| COMPLETE FOR HEPATITIS B, HEPATITIS C, AND HIV ONLY | | | |
| STAGING | HEPATITIS B | HEPATITIS C | HIV |
| ACUTE | <input type="checkbox"/> | <input type="checkbox"/> | N/A |
| CHRONIC | <input type="checkbox"/> | <input type="checkbox"/> | N/A |
| NEW DIAGNOSIS | N/A | N/A | <input type="checkbox"/> |
| OLD CASE - PREVIOUSLY DIAGNOSED/KNOWN IN MB | <input type="checkbox"/> | <input type="checkbox"/> | <input type="checkbox"/> |
| PREVIOUS DIAGNOSIS – NEW TO MANITOBA | <input type="checkbox"/> | <input type="checkbox"/> | <input type="checkbox"/> |
| UNKNOWN/UNDETERMINED | <input type="checkbox"/> | <input type="checkbox"/> | N/A |
| COMPLETE FOR SYPHILIS ONLY | | | |
| SYPHILIS SIGNS/SYMPTOMS (CHECK ALL THAT APPLY) | | | |
| <input type="checkbox"/> ASYMPTOMATIC | <input type="checkbox"/> GENITAL ULCER | <input type="checkbox"/> OCULAR INVOLVEMENT | <input type="checkbox"/> OTHER (SPECIFY): |
| <input type="checkbox"/> ANAL ULCERATIVE LESIONS | <input type="checkbox"/> HAIR LOSS (ALOPECIA) | <input type="checkbox"/> ORAL ULCERATIVE LESIONS | |
| <input type="checkbox"/> CONDYLOMATA LATA | <input type="checkbox"/> MENINGITIS | <input type="checkbox"/> RASH | |
| <input type="checkbox"/> PRIMARY | <input type="checkbox"/> LATE LATENT (GREATER THAN 1 YEAR AFTER INFECTION) | | <input type="checkbox"/> UNKNOWN/UNDETERMINED |
| SYPHILIS STAGING <input type="checkbox"/> SECONDARY | <input type="checkbox"/> TERTIARY | | <input type="checkbox"/> OLD CASE-PREVIOUSLY DIAGNOSED/ KNOWN IN MB |
| <input type="checkbox"/> EARLY LATENT (LESS THAN 1 YEAR AFTER INFECTION) | | | |
| ADDITIONAL PRESENTATIONS (SITES) | <input type="checkbox"/> NEUROSYPHILIS | <input type="checkbox"/> GUMMATOUS SYPHILIS | <input type="checkbox"/> CARDIOVASCULAR SYPHILIS |

VI. RISK FACTOR INFORMATION (OPTIONAL)

[subject](#) > [risk factors](#)

| |
|---|
| SPECIFY KNOWN APPLICABLE RISK FACTORS IF RELEVANT TO THE PUBLIC HEALTH INVESTIGATION: |
| |

VII. ADDITIONAL INFORMATION (IF APPLICABLE)

| | |
|--|--|
| ADDITIONAL UPDATES OR INFORMATION | |
| FOR LABORATORY CONFIRMED CASES ONLY | |
| HAS CLIENT BEEN INFORMED OF DIAGNOSIS? <input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> PENDING | |
| (HIV CASE ONLY) HAS NEED FOR HIV DISCLOSURE WITH PARTNERS BEEN DISCUSSED? <input type="checkbox"/> YES <input type="checkbox"/> NO | |

VIII. REPORTER INFORMATION

| | |
|-----------------------------------|---------------------------------|
| FORM COMPLETED BY (PRINT NAME) | FACILITY NAME/ ADDRESS/ PHONE # |
| SIGNATURE | |
| FORM COMPLETION DATE (YYYY-MM-DD) | |

REMINDER: TESTING FOR ALL STBBI IS RECOMMENDED

| | |
|-------------------|---------------------------|
| CLIENT LAST NAME: | CLIENT FIRST NAME: |
| PHIN: _____ | OR DOB (YYYY-MM-DD) _____ |

IX. CONTACTS OF CASE (FOR LAB CONFIRMED CASES ONLY)

investigation quick entry > exposure summary > create
transmission event > known contacts
contact investigation > disposition / intervention

COPY PAGE IF REQUIRED. PLEASE PROVIDE AS MUCH DETAIL AS POSSIBLE.

CASE DECLINED TO IDENTIFY CONTACTS NUMBER OF ANONYMOUS CONTACTS _____

| CONTACT PERSONAL INFORMATION | PREGNANT? | WHO WILL NOTIFY? | EXPOSURE START AND END DATES YYYY-MM-DD |
|---|--------------------------|--|---|
| NAME: PHIN (IF KNOWN): DOB/AGE: ADDRESS: PHONE: ALTERNATE CONTACT INFO (E.G PHONE, SOCIAL MEDIA, EMAIL): | <input type="checkbox"/> | <input type="checkbox"/> PUBLIC HEALTH <input type="checkbox"/> CASE <input type="checkbox"/> HEALTH CARE PROVIDER | START DATE END DATE |
| NAME: PHIN (IF KNOWN): DOB/AGE: ADDRESS: PHONE: ALTERNATE CONTACT INFO (E.G PHONE, SOCIAL MEDIA, EMAIL): | <input type="checkbox"/> | <input type="checkbox"/> PUBLIC HEALTH <input type="checkbox"/> CASE <input type="checkbox"/> HEALTH CARE PROVIDER | START DATE END DATE |
| NAME: PHIN (IF KNOWN): DOB/AGE: ADDRESS: PHONE: ALTERNATE CONTACT INFO (E.G PHONE, SOCIAL MEDIA, EMAIL): | <input type="checkbox"/> | <input type="checkbox"/> PUBLIC HEALTH <input type="checkbox"/> CASE <input type="checkbox"/> HEALTH CARE PROVIDER | START DATE END DATE |
| NAME: PHIN (IF KNOWN): DOB/AGE: ADDRESS: PHONE: ALTERNATE CONTACT INFO (E.G PHONE, SOCIAL MEDIA, EMAIL): | <input type="checkbox"/> | <input type="checkbox"/> PUBLIC HEALTH <input type="checkbox"/> CASE <input type="checkbox"/> HEALTH CARE PROVIDER | START DATE END DATE |

PLEASE SUBMIT THIS FORM BY SECURED FAX OR COURIER TO THE MANITOBA HEALTH SURVEILLANCE UNIT.
4050 – 300 CARLTON ST. WINNIPEG, MB | CONFIDENTIAL FAX 204-948-3044
AFTER HOURS EMERGENCY PHONE FOR PUBLIC HEALTH ISSUES IS (204) 788-8666.

THIS FORM (AND GUIDANCE FOR COMPLETION) IS AVAILABLE FOR DOWNLOAD IN A FILLABLE PDF FORMAT AT:
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