Infectious and congenital syphilis in Canada, 2010–2015

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Abstract

Background: Syphilis is the third most commonly reported notifiable sexually transmitted infection (STI) in Canada, following chlamydia and gonorrhea, respectively. Rates of this STI have been rising rapidly in Canada since 2001.

Objective: To summarize trends observed in syphilis rates for 2010 to 2015 in Canada.

Methods: Laboratory-confirmed cases of infectious syphilis and early congenital syphilis were reported to the Public Health Agency of Canada by all of the Canadian provinces and territories. National infectious syphilis rates were computed, as were rates per sex, age group and province/territory. Rates of congenital syphilis were also calculated.

Results: From 2010 to 2015, the rate of infectious syphilis in Canada increased by 85.6%, from 5.0 to 9.3 cases per 100,000 population. In 2015, a total of 3,321 cases of infectious syphilis were reported, mainly in males (93.7%), among whom the rate was 17.5 cases per 100,000 males versus 1.2 per 100,000 females. The rate also rose faster among males in 2010–2015, a 90.2% increase versus 27.8% among females. Individuals aged 20–39 years had the highest rates. Across the provinces and territories, the highest rates of infectious syphilis were in Nunavut, British Columbia and Manitoba.

The rate of congenital syphilis decreased from 2010 to 2014 (1.6 to 0.3 cases per 100,000 live births) before increasing to 1.5 cases per 100,000 live births in 2015, which corresponds to six reported cases.

Conclusion: Rates of syphilis continue to rise in Canada, especially among young men, and this is consistent with trends in the United States of America and European Union. Based on data from Canada and from these regions, the sexual behaviour of men who have sex with men (MSM) is thought to be a major risk factor for syphilis.


Introduction

Syphilis, caused by the bacterium Treponema pallidum (1), is the third most commonly reported notifiable sexually transmitted infection (STI) in Canada, after chlamydia and gonorrhea, respectively. If left untreated, a primary syphilis infection can progress through secondary, latent and tertiary disease stages (2). Out of the four stages of syphilis, only three are infectious and therefore of public health significance: primary, secondary and early latent syphilis (1). Neurological symptoms can occur at any stage. Symptoms of infection in the earlier stages include chancres, condyloma lata and generalized lymphadenopathy. Cardiologic and musculoskeletal manifestations may occur if the infection remains untreated and reaches the tertiary stage. There is a synergy between HIV and syphilis as syphilis infection increases HIV viral load and HIV transmission (2). Moreover, for exposed individuals, HIV acquisition is two- to five-fold higher among those infected with syphilis than those without syphilis infection (2).

Congenital syphilis occurs through mother-to-child transmission, mainly in utero but also at birth. Congenital syphilis may have severe consequences for the newborn, such as cerebral palsy, hydrocephalus, sensorineural hearing loss, musculoskeletal deformity or death (3). The risk of transmission varies from 10% to more than 70% depending on the mother’s stage of disease (3). Transmission may be prevented with timely diagnosis and adequate treatment.

After years of low incidence among both males and females, there has recently been a large increase in the number of syphilis cases, mainly among males (1). This rate increase coincides with the growing number of outbreaks reported in several cities and provinces across Canada among men who have sex with men (MSM), and especially among HIV-infected MSM, the heterosexual population and some Indigenous communities (1).
The objective of this article is to summarize observed trends in reported laboratory-confirmed infectious syphilis and congenital syphilis rates in Canada in the period 2010 to 2015. Rates were analyzed by sex, age and geographic distribution.

Methods

Data sources

Provincial and territorial health authorities provide non-nominal data on laboratory-confirmed cases to the Public Health Agency of Canada (PHAC) through the Canadian Notifiable Disease Surveillance System (CNDSS) (4). Confirmed case definitions of infectious syphilis and congenital syphilis are presented in the Appendix (5).

Variables submitted along with the diagnosis include sex, age at time of diagnosis, year of diagnosis and province/territory of diagnosis. All stages are notifiable but only infectious stages (primary, secondary and early latent) were included in this report. The received data were validated in collaboration with the corresponding province or territory. Data from January 1, 2010 to December 31, 2015 were available from all provinces and territories and were extracted from the CNDSS in July 2017.

Data analysis

Descriptive analysis was performed using Microsoft Excel. National annual reported case rates of infectious syphilis were computed per 100,000 population (or per males or females) for all years using number of cases from the CNDSS as numerators, and Statistics Canada yearly population estimates as denominators. Sex, age group and province/territory-specific rates were also calculated. For congenital syphilis, rates were computed per 100,000 live births. For 2014 and 2015, preliminary numbers of live births drawn from Statistics Canada were used, as final numbers were not yet available. No statistical procedures were used for comparative analyses. Small numbers are more susceptible to change and so corresponding rates should be interpreted with caution. Previous reports may provide different rates for some years due to reporting delays and data updating.

Results

From 2010 to 2015, the rate of reported laboratory-confirmed cases of infectious syphilis in Canada increased from 5.0 to 9.3 cases per 100,000 population (Figure 1). This represents an increase of 85.6% over this time period.

Sex and age

In 2015, a total of 3,321 cases of infectious syphilis were reported, of which 93.7% were males. This was reflected in the much higher rate of infectious syphilis among males than among females (17.5 cases per 100,000 males versus 1.2 cases per 100,000 females). The rate of infectious syphilis also increased faster among males than among females in 2010–2015 (90.2% among males versus 27.8% among females) (Figure 1).

Geographic distribution

The three provinces with the highest reported rates of laboratory-confirmed syphilis in 2015 were Nunavut, British Columbia and Manitoba (Table 1). The greatest rate decreases were seen in the Northwest Territories, Saskatchewan and New Brunswick.
Table 1: Number and rate of reported laboratory-confirmed cases of infectious syphilis, by province and territory in Canada, 2010–2015

<table>
<thead>
<tr>
<th>Province or territory</th>
<th>Laboratory-confirmed cases by year of diagnosis (rate per 100,000)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>2010</td>
</tr>
<tr>
<td>Alberta</td>
<td>173</td>
</tr>
<tr>
<td>British Columbia</td>
<td>92</td>
</tr>
<tr>
<td>Manitoba</td>
<td>17</td>
</tr>
<tr>
<td>New Brunswick</td>
<td>34</td>
</tr>
<tr>
<td>Newfoundland and Labrador</td>
<td>4</td>
</tr>
<tr>
<td>Northwest Territories</td>
<td>3</td>
</tr>
<tr>
<td>Nova Scotia</td>
<td>18</td>
</tr>
<tr>
<td>Nunavut</td>
<td>0</td>
</tr>
<tr>
<td>Ontario</td>
<td>774</td>
</tr>
<tr>
<td>Prince Edward Island</td>
<td>0</td>
</tr>
<tr>
<td>Quebec</td>
<td>546</td>
</tr>
<tr>
<td>Saskatchewan</td>
<td>36</td>
</tr>
<tr>
<td>Yukon</td>
<td>0</td>
</tr>
<tr>
<td>Canada</td>
<td>1,697</td>
</tr>
</tbody>
</table>

Congenital syphilis
The number of laboratory-confirmed cases of congenital syphilis reported in Canada varied from one to six cases per year in 2010–2015 (Table 2).

Table 2: Number of reported laboratory-confirmed cases and rates of congenital syphilis, 2010–2015, Canada

<table>
<thead>
<tr>
<th>Indicator</th>
<th>Year of diagnosis</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>2010</td>
</tr>
<tr>
<td>Number of cases</td>
<td>6</td>
</tr>
<tr>
<td>Rate per 100,000 live births</td>
<td>1.59</td>
</tr>
</tbody>
</table>

Discussion
In Canada, the rates of reported cases of infectious syphilis markedly increased from 2010 to 2015. The burden of syphilis seems to be disproportionately placed on men. Other high-income countries such as the United States of America (USA), Australia and the United Kingdom have reported similar increases in numbers of cases and rates (6-8).

The very high number of cases among males is thought to be mainly because of an increase in cases among MSM (1). Newfoundland and Labrador and Manitoba are among the provinces with the highest increases over 2010–2015. These provinces have reported that increased diagnoses among MSM was the main factor driving rates upwards (9,10). This is of concern as syphilis contraction increases the probability of acquiring and transmitting HIV (1). A greater increase of reported syphilis cases has been observed among MSM living with HIV than among noninfected MSM in both Western Europe and the USA (11).

The causes of the increase in the rate of syphilis among MSM are multifactorial and complex. Changing community norms and behaviours as well as new preventive interventions such as pre-exposure prophylaxis (PrEP) might explain this rise. Many MSM have adopted behaviour patterns such as serosorting (choosing to have condomless sex with partners with the same HIV status) or having condomless oral sex, with the intention of decreasing HIV transmission (12,13). However, the lowered condom use might increase the risk of contracting other STIs (13,14). In a Toronto-based study, condomless anal sex with casual partners in the previous six months was associated with syphilis infection among MSM living with HIV (15). Also of concern is the fact that Internet-based social media are increasingly being used to easily find sex partners. This may promote concurrent partnerships and rates of acquisition of new partners and decrease intervals between sex partners (16). The social mixing patterns with the use of saunas, and the consumption of recreational drugs that may impair judgment in making decisions about sexual acts are also risk factors for acquisition and transmission of syphilis and other STIs (16,17). Lastly, increased risk-taking behaviours and a rise in STI incidence have been reported among HIV-negative MSM using PrEP (18-20). In Canada, PrEP was not significantly used in 2010–2015. However, data from other countries and from research studies highlight the importance of frequent STI screening of MSM on PrEP to ensure that symptomatic and asymptomatic STIs are treated in a timely way to halt transmission, as PrEP use increases over time (21).

This finding highlights the importance of public health action to mitigate transmission of syphilis and identify new risk groups, such as MSM on PrEP.

The rate increase in women is also worrying as congenital syphilis tends to increase with rates of primary and secondary syphilis among women of childbearing age. A recent study on the epidemiology of syphilis in Winnipeg reported that one quarter...
of women with syphilis were pregnant at diagnosis (22). No cases of congenital syphilis were found in the study.

This finding highlights the importance of universal screening of pregnant women in a context of syphilis resurgence in Canada, as recommended by the Canadian Guidelines on Sexually Transmitted Infections (1).

**Strengths and limitations**

This surveillance report presents a national portrait of the current infectious syphilis epidemiology and was based on data from all provinces and territories. It describes sex, age and province/territory-specific rates over a six-year period.

Some limitations of the data should be noted. First, some numbers of cases of infectious and congenital syphilis were low. This leads to less stable rates, especially for congenital syphilis for which less than 10 cases were reported annually in Canada in 2010–2015. Therefore, variations in rates over time should be interpreted with caution. Second, these figures likely underestimate the incidence rate of syphilis from 2010 to 2015 as some infections may be asymptomatic, unscreened, undiagnosed or unreported. Screening, laboratory testing and reporting practices are heterogeneous across provinces and territories, and reports to the PHAC of syphilis cases by stage vary between provinces/territories. Therefore, we were not able to calculate valid stage-specific rates or to report on the number of cases of specific conditions such as neurosyphilis. Likewise, although age structures may vary across provinces and territories, we did not perform standardization by age. Therefore, direct comparison between provinces should be made with caution.

Trend analysis on the data was not performed, which is a limitation of this report. Lastly, risk factors and clinical presentation are not available in this surveillance system, preventing identifying risk factors associated with the observed increased rates.

**Conclusion**

In conclusion, syphilis rates in Canada have risen markedly over time. Males make up the vast majority of syphilis cases, and based on data from Canada, USA, Australia and other countries, MSM are one of the groups at highest risk. A better understanding of transmission dynamics and social and sexual networking is needed to guide prevention efforts.

**Authors’ statement**

YC – Conceptualization, methodology, writing – original draft
JM – Software, data collection and curation, validation, formal analysis, visualization, writing – review and editing
JS – Writing – original draft, visualization
AL – Validation, formal analysis, visualization, writing – review and editing
JA – Conceptualization, writing – original draft

**Conflict of interest**

None.

**Contributors**

Chris Archibald: Supervision, writing – review and editing, resources, project administration
Jennifer Siushansian: writing – review and editing

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**References**


### Appendix: Case definitions of confirmed cases of infectious and congenital syphilis

**Laboratory confirmation of early congenital syphilis infection (within two years of birth):**
- Identification of *Treponema pallidum* by dark-field microscopy, fluorescent antibody or equivalent examination of material from nasal discharges, skin lesions, placenta, umbilical cord or autopsy material of a neonate (up to four weeks of age)
  OR
- Reactive serology (non-treponemal and treponemal) from venous blood (not cord blood) in an infant/child with clinical, laboratory or radiographic evidence of congenital syphilis whose mother is without documented evidence of adequate treatment
  OR
- Detection of *T. pallidum* DNA in an appropriate clinical specimen

**Laboratory confirmation of primary syphilis infection:**
- Identification of *T. pallidum* by dark-field microscopy, fluorescent antibody, nucleic acid testing or equivalent examination of material from a chancre or a regional lymph node
  OR
- Presence of one or more typical lesions (chancres) and reactive treponemal serology, regardless of non-treponemal test reactivity, in individuals with no previous history of syphilis
  OR
- Presence of one or more typical lesions (chancres) and a fourfold or greater increase in the titre over the last known non-treponemal test in individuals with a past history of syphilis treatment

**Laboratory evidence of infection for secondary syphilis:**
- Identification of *T. pallidum* by dark-field microscopy, fluorescent antibody, nucleic acid testing or equivalent examination of mucocutaneous lesions, condylomata lata and reactive serology (non-treponemal and treponemal)
  OR
- Presence of typical signs or symptoms of secondary syphilis (e.g. mucocutaneous lesions, alopecia, loss of eyelashes and lateral third of eyebrows, iritis, generalized lymphadenopathy, fever, malaise or splenomegaly) AND either a reactive serology (non-treponemal and treponemal) OR a fourfold or greater increase in titre over the previous known non-treponemal test

**Laboratory confirmation of early latent syphilis infection (<1 year after infection):**
An asymptomatic patient with reactive serology (treponemal and/or non-treponemal) who, within the previous 12 months, had one of the following:
- Nonreactive serology
- Symptoms suggestive of primary or secondary syphilis
- Exposure to a sexual partner with primary, secondary or early latent syphilis

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*Quebec's definition requires the use of two tests including a treponemal one for a diagnostic of primary, secondary or early latent syphilis*