A cost-benefit/cost-effectiveness analysis of an unsanctioned supervised smoking facility in the Downtown Eastside of Vancouver, Canada

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Abstract

Background: Smoking crack involves the risk of transmitting diseases such as HIV and hepatitis C (HCV). The current study determines whether the formerly unsanctioned supervised smoking facility (SSF)—operated by the grassroots organization, Vancouver Area Network of Drug Users (VANDU) for the last few years—costs less than the costs incurred for health-care services as a direct consequence of not having such a program in Vancouver, Canada.

Methods: The data pertaining to the attendance at the SSF was gathered in 2012–2013 by VANDU. By relying on this data, a mathematical model was employed to estimate the number of HCV infections prevented by the former facility in Vancouver’s Downtown Eastside (DTES).

Results: The DTES SSF’s benefit-cost ratio was conservatively estimated at 12.1:1 due to its low operating cost. The study used 70% and 90% initial pipe-sharing rates for sensitivity analysis. At 80% sharing rate, the marginal HCV cases prevented were determined to be 55 cases. Moreover, at 80% sharing rate, the marginal cost-effectiveness ratio ranges from $1,705 to $97,203. The results from both the baseline and sensitivity analysis demonstrated that the establishment of the SSF by VANDU on average had annually saved CAD$1.8 million dollars in taxpayer’s money.

Conclusions: Funding SSFs in Vancouver is an efficient and effective use of financial resources in the public health domain; therefore, Vancouver Coastal Health should actively participate in their establishment in order to reduce HCV and other blood-borne infections such as HIV within the non-injecting drug users.

Keywords: Supervised smoking facility, Crack, VANDU, Hepatitis C, Downtown Eastside

Background

Smoking crack cocaine is not only on the rise in the Canadian municipalities, but it is also often neglected by health officials—especially so when compared to similar inner-city health problems such as injection drug use [1–4]. In British Columbia, the daily usage of crack cocaine within the general population is higher than that in any other provinces within Canada [5]. This is a pressing problem in Vancouver, where daily crack use, within a cohort of injection drug users, increased from 7.4% in 1996 to 42.6% in 2005 [6]. Among drug users in Vancouver’s Downtown Eastside (DTES), the rate of crack use has been reported to be as high as 86.6% [7]. The use of crack is associated with several other risks when compared to the tendencies displayed by other drug-using populations. For example, crack users are more likely to have unstable housing [8], be involved in sex work [9], participate in risky behavior [10–12], engage in criminal activity [13,14], experience multiple health problems [5], and are less likely to access social and health services [15].

Research conducted upon a cohort of crack-user population in Vancouver’s DTES revealed that participants had reported 80% sharing rate as it is related to their drug smoking paraphernalia [16,17]. Studies have shown a higher-than-average prevalence of human immunodeficiency virus (HIV), hepatitis C virus (HCV), and tuberculosis in users of crack cocaine who report no injection drug use [17]. However, the evidence of the relationship between non-injecting drug use and HIV/HCV infection is ambiguous. Some researchers have suggested
that non-injecting drug users (NIDUs) are often involved in unsafe sexual behavior [18] and that HCV transmission in NIDUs is associated with tattooing [19]. Some researchers have stirred up a controversy in suggesting that NIDUs are essentially injecting drug users (IDUs) who have failed to report their route of transmission accurately [20].

Nevertheless, research conducted on NIDUs suggests that infectious diseases may have been transmitted by the sharing of crack pipes [21,22]. In fact, most users are oblivious to the risks involved in sharing drug tools [21]. Some researchers postulate that HIV and HCV transmission can be accounted for by the high prevalence of oral lesions in crack smokers. Some of these include sores, blisters, and cuts on their lips and oral cavities—caused because of the mouth and lips coming in contact with hot glass, hot smoke, and the sharp edges of glass pipe stems or metal pipe stems [21]. The lack of knowledge with respect to transmittable diseases further engenders and reinforces the reckless exchange of drug equipment. In fact, a study demonstrated that 2% of crack pipes tested positive for HCV [23].

Scientific evaluation of Insite, North America’s first and only supervised injection facility, showed that it has successfully reduced needle sharing and overdose death while concurrently improving service uptake and public order within the DTES [24-26]. Despite the improvement of conditions in the DTES after the opening of Insite, Vancouver is still riddled with concerns regarding public health and order related to drug use, including crack and crystal methamphetamine [27,28]. Accordingly, the region’s health authority has shown some interest in applying for an exemption under the Controlled and Substance Act of the Criminal Code of Canada to open a supervised smoking facility (SSF) in the DTES. However, the concept of a government-sanctioned SSF is somewhat controversial, particularly because the potential impact and benefits of such a facility are unknown.

Therefore, the present research was conducted to determine whether a case could be made for the establishment of SSFs in the DTES of Vancouver. Specifically, the current study analyzed the cost-benefits and cost-effectiveness of the only SSF in Canada, operated by Vancouver Area Network of Drug Users (VANDU) without a license for a few years. The SSF mentioned above was located in VANDU’s front office in the DTES, along East Hastings Street. VANDU has over 800 volunteers, 1,300 active members [29], and a Board of Directors composed of current and former users. See Figure 1 for the location of VANDU in the DTES.

In December of 2013, VANDU was forced to shut down the SSF under the direction of their funding agency, the Vancouver Coastal Health. Using mathematical modelling with conservative parameter estimates, this analysis estimated the number of HCV infections prevented as a result of SSF. The savings from illnesses avoided were compared to the operational cost of a SSF. The analysis was eventually extended to consider the impact of opening additional SSFs in the DTES.

**Methods**

**Background**

VANDU operates on an annual budget of CAD$200,000 funded through Vancouver Coastal Health. One of their various programs included the operation of an unsanctioned SSF. The smoking room was operated by peers and was accessible to one person at a time. There would be an unusually big lineup to use the room that contained a fan. Within VANDU, NIDUs would generally be provided with a ‘safer crack use kit’ that contained the following: mouth pieces, wooden push sticks, screens, alcohol swaps, and heat-resistant and shatter-proof glass pipes which minimized chances of injury to the users’ lips and mouth. See Figure 2 for materials contained in the ‘safer crack use kit’ provided at VANDU.

Moreover, VANDU’s SSF provided a clean and safe environment within which one could use pre-obtained illicit drugs, get medical attention in the event of an overdose, and obtain access or referral to primary health care when required. This study was approved by Simon Fraser University Research Ethics Board (study number: 2013 s0058). VANDU’s Executive Board also approved the study since it corresponded with its philosophy and the demand that all projects directly involve its members.

**Model**

For this analysis, it was necessary to calculate the effects of both providing clean equipment as well as that of adopting safer smoking behaviors. Along the lines of research conducted on the economic impact of a needle exchange program in Edmonton, Alberta, Canada, this study uses a mathematical model to estimate the number of HCV infections that could be prevented through the establishment of a SSF [30]. The number of new HCV infections avoided, $A$, is calculated as follows:

$$A = INs(1−(1−qt)^m),$$

where $m$ is the number of sharing partners when pipes are shared, $t$ is the probability of HCV transmission when using an HCV-infected pipe, $s$ is the rate of pipe sharing, $I$ is the proportion of NIDU population that is HCV negative, $N$ is the number of pipes in circulation, $d$ is the percentage of pipes not cleaned before use, and $q$ is the proportion of the NIDU population that is HCV positive.

Initially, this study was meant to use a few other mathematical models such as those of Kaplan and O’Keefe [31], Lurie and Drucker [32], Gold et al. [33], Laufer [34], and
Pinkerton [35,36]. However, due to lack of data, such as the rate of HIV transmission from a single pipe and the rate of secondary transmission, this analysis had to rely on the Jacobs et al. [30] model. However, the model employed in the current study has previously been adopted by four different studies [37-40], which have found that this model is the best choice for predicting actual and potential cases of HIV and HCV in a Canadian setting. Moreover, the model employed in this study has successfully produced estimates of HIV and HCV cases within the IDU population—similar to known data widely cited in peer-reviewed reports.

Additionally, this study also uses behavior change incorporated by previous costing studies conducted on supervised injection facilities [37-41] because of the empirical evidence it provides [24,35,41,42]. Although previous costing studies often go wrong when it comes to using caution and employing an odd ratio of 0.60, this study uses the point estimate of 0.30 used by previous studies [37-42] and estimated by Kerr et al. [24]. The data collected by VANDU during 2012–2013 pertains to the number of visits per month to the SSF.

**Variables and parameters**

Medical and scientific literatures were used in cases when Vancouver-specific data was not available. Where estimates differed, this study used the lower bound, so all estimates remain conservative. The concept of behavioral change in
the NIDU population was adopted based on the behavioral changes related to IDU needle-sharing behavior outside of the SIF in Vancouver. Kerr et al. [24] and Bravo et al. [43] found that IDUs who relied upon SIFs were also able to reduce their needle-sharing activities outside of the facility up to a significant extent. Table 1 provides the estimates and variables used in the model (please note that percentages need to be converted to fractions when inputting the variables in the model).

Consequently, it was presumed that NIDUs that visited the unsanctioned SSF were less likely to share their pipes with others outside of the facility. Furthermore, if a second SFF was established, the behavioral change on pipe sharing would occur only if new NIDUs became users of SSF [28]. On the contrary, if the SSF was frequented by the current users, thereby restricting its use to current users simply indulging in some additional smoking, no further behavioral changes can be assumed. Accordingly, behavioral change is only accounted for in the second facility.

Furthermore, since there was no estimated number of crack users in the DTES of Vancouver, this number was calculated based on the percentage of drug users (conservatively estimated to be around 5,000 in the DTES [51-53]) who have smoked crack. The total number of drug users was reported by DeBeck et al. [7] to be 86.6% (5,000 drug users × 0.866 use crack = 4,330 crack users in the DTES). This number was subsequently multiplied by the number of subjects who smoked crack per day—estimated to be around ten per day [54,55] (4,330 × 10 × 365 days = 15,804,500 smoking per year).

The number of those indulging in smoking per year was multiplied by the percentage of pipe sharing in the Downtown Eastside (15,804,500 smoking per year × 0.80 sharing = 12,643,600 shared crack smoking events). The total visits to the SSF during 2012–2013 year were determined to be 23,120 per year with the average visit of 1,843 per month. Consequently, 17,696 smoking incidents were not shared as a result of having an unsanctioned SSF operating in the DTES (23,120 × 0.8 sharing = 17,696 smoking events that were not shared). This number was added to the behavioral change odd ratio and later deducted from the total shared crack pipe events in the DTES.

**The medical cost of new HCV cases**

HCV infection among people who use drugs is a serious pressing concern in Canada and the United States [56,57]. HCV infections could lead to multiple health problems such as cirrhosis, liver failure, hepatocellular carcinoma, and even death [58]. Accordingly, 50% of patients achieve sustained virological success to treatment [59]. Pegylated interferon, in combination with ribavirin, is the standard course of treatment for HCV-infected patients [57]. The range of treatment for HCV patients is determined based on the genotype: ‘a 48-week course is recommended for genotypes 1 and 4, whereas a 24-week course is recommended for genotypes 2 and 3’ [57], p. 1016. Accordingly, the cost of treatment varies according to genotype and seriousness of infections.

On an average, savings from HCV range from $20,000 per completed course of treatment per patient [60], to $30,000 [61], and to more than $69,188 [58]. This study uses a conservative figure of CAD$35,143 (2012 US Dollars = 33,856), as reported in [62] and utilized in costing studies of a potential SIF in Montreal [39] and Ottawa [40]. The conservative figure used in this study essentially disregards the cost of the complications arising from HCV in hepatocellular carcinoma, liver failure, and liver transplant cases.

**Cost of SSF**

In order to estimate the cost of operating a potential SSF, it was important to calculate the operating cost of

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**Table 1 Sources for variables used in mathematical modelling**

| Variable                                      | Value | Source                      |
|-----------------------------------------------|-------|-----------------------------|
| Rate of pipe sharing (s)                      | 80.0% | Ivins et al. [44]           |
| Number of pipes in circulation (N)            | 90,000| VANDU [45], Mui [46]        |
| Percentage of pipes not cleaned (d)           | 33.0% | Scheinmann et al. [47]      |
| Number of sharing partners (m)                | 6.30  | Gyarmathy and Neaigu [48]   |
| Proportion of crack users who are HCV negative (I) | 83.0% | Fischer et al. [49]        |
| Proportion of crack users who are HCV positive (q) | 17.0% | Fischer et al. [49]        |
| Probability of HCV infection from single crack pipe (t) | 2.00% | Fischer et al. [23], Gilbert et al. [50] |

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**Figure 2** Content materials of a ‘safer crack use kit’ provided at VANDU.
the existing SSF in the DTES. The former facility operated from Monday to Friday from 10–7 pm. On weekends, the facility would operate from 4–7 pm. The staff supervising the unsanctioned SSF were mostly volunteers that were provided with a small stipend, collectively amounting to CAD$47,203 per year. The total cost of the rent and the safe crack kit is estimated to be CAD$50,000. Altogether, the operating cost of the facility is estimated to be CAD$97,203.

Results
The model used here [26] predicted the number of new HCV cases prevented based on the pipe-sharing rate. This included the impact of behavioral changes in pipe sharing outside of the SSF. The behavioral change, according to Tables 2 and 3, was only considered twice—once for the first SSF and once for the second SSF—based on a conservative odd ratio that falls within the limit specified by [31]. As expected, the results presented in Tables 2 and 3 show that expanding SSFs would decrease HCV cases. The model predicts 57–114 cases for HCV with the marginal range being much smaller at 1–57 for HCV.

This range disparity, as outlined in Tables 2 and 3, translates into substantial differences between the cumulative estimates and the marginal estimates. For example, according to Table 2, the benefit-cost ratio ranges from 5.9 to 20.6 and the cost-effectiveness value ranges from $1,705 to $5,969 (cost per lifetime treatment).

### Table 2 The cumulative cost-effectiveness and cost-benefit of SSF in Vancouver using Jacobs et al.’s [30] model

| Variables | Annual cost of operation ($) | Sharing rate (%) | # of HCV averted | Cost-effectiveness ratio HCV ($) | Cost-benefit ratio HCV |
|-----------|-------------------------------|------------------|-----------------|---------------------------------|-----------------------|
| Post SSF  | 97,203                        | 69               | 57              | 1,705                           | 20.6                  |
|           |                               | (78, 60)         | (65, 50)        | (1,495, 1,944)                  | (23.5, 18.1)          |
| Two SSF   | 194,406                       | 59               | 109             | 1,784                           | 19.7                  |
|           |                               | (67, 52)         | (121, 93)       | (1,607, 2,090)                  | (21.9, 16.8)          |
| Three SSF | 291,609                       | 58               | 110             | 2,651                           | 13.3                  |
|           |                               | (67, 52)         | (121, 94)       | (2,410, 3,102)                  | (14.6, 11.3)          |
| Four SSF  | 388,812                       | 58               | 111             | 3,503                           | 10                    |
|           |                               | (67, 52)         | (122, 94)       | (3,187, 4,136)                  | (11, 8.5)             |
| Five SSF  | 486,015                       | 58               | 112             | 4,339                           | 8.1                   |
|           |                               | (67, 52)         | (123, 95)       | (3,951, 5,116)                  | (8.9, 6.9)            |
| Six SSF   | 583,218                       | 57               | 113             | 5,161                           | 6.8                   |
|           |                               | (66, 52)         | (124, 95)       | (4,703, 6,139)                  | (7.5, 5.7)            |
| Seven SSF | 680,421                       | 57               | 114             | 5,969                           | 5.9                   |
|           |                               | (66, 52)         | (124, 96)       | (5,487, 7,088)                  | (6.4, 5)              |

The numbers in parentheses represent the results of the sensitivity analysis (90% sharing rate, 70% sharing rate).

### Table 3 The marginal cost-effectiveness and cost-benefit of SSF in Vancouver using Jacobs et al.’s [30] model

| Variables | Annual cost of operation ($) | Sharing rate (%) | # of HCV averted | Cost-effectiveness ratio HCV ($) | Cost-benefit ratio HCV |
|-----------|-------------------------------|------------------|-----------------|---------------------------------|-----------------------|
| Post SSF  | 97,203                        | 69               | 57              | 1,705                           | 20.6                  |
|           |                               | (78, 60)         | (65, 50)        | (1,495, 1,944)                  | (23.5, 18.1)          |
| Two SSF   | 97,203                        | 59               | 52              | 3,739                           | 18.8                  |
|           |                               | (67, 52)         | (56, 43)        | (1,736, 2,261)                  | (20.2, 15.5)          |
| Three SSF | 97,203                        | 58               | 1               | 97,203                          | 0.4                   |
|           |                               | (67, 52)         | (1, 1)          | (97,203, 97,203)                | (0.4, 0.4)            |
| Four SSF  | 97,203                        | 58               | 1               | 97,203                          | 0.4                   |
|           |                               | (67, 52)         | (1, 0.5)        | (97,203, 194,406)               | (0.4, 0.2)            |
| Five SSF  | 97,203                        | 58               | 1               | 97,203                          | 0.4                   |
|           |                               | (67, 52)         | (1, 0.5)        | (97,203, 194,406)               | (0.4, 0.2)            |
| Six SSF   | 97,203                        | 57               | 1               | 97,203                          | 0.4                   |
|           |                               | (66, 52)         | (1, 0.5)        | (97,203, 194,406)               | (0.4, 0.2)            |
| Seven SSF | 97,203                        | 57               | 1               | 97,203                          | 0.4                   |
|           |                               | (66, 52)         | (1, 0.5)        | (97,203, 194,406)               | (0.4, 0.2)            |

The numbers in parentheses represent the results of the sensitivity analysis (90% sharing rate, 70% sharing rate).
In contrast, the marginal estimates of SSF expansion translate into a much smaller return. This is particularly true with respect to its benefit-cost and cost-effectiveness ratio; for instance, the marginal benefit-cost ratio varies from 20.6 to 0.4. The marginal cost-effectiveness value for HCV ranges from $1,705 to $97,203 (cost per lifetime treatment). Furthermore, Table 3 shows that both cumulative benefit-cost and cost-effectiveness ratios dwindle after the second SSF.

Finally, a sensitivity analysis was conducted for the models employed. The sensitivity analysis pertained to simulating different pipe-sharing rates (see Tables 2 and 3). Similar to costing studies in Vancouver [37,38], Montreal [39], and Ottawa [40] that used different needle-sharing rates, the current analysis used 70% and 90% initial pipe-sharing rates. Convincingly, the results from both the baseline and sensitivity analysis demonstrate that the establishment of an SSF by VANDU had saved taxpayer money.

Discussion
The current analyses assessed whether the former SSF, operated by VANDU in the DTES, would have had a net positive fiscal impact on the Canadian society and whether or not this policy initiative would save public health-care funds by averting new HCV infections. Moreover, the optimal number of SSFs was assessed based on marginal cost-effectiveness and benefit-to-cost ratios. The results presented here suggest that closing the only unsanctioned SSF in Vancouver was a policy failure that has potentially resulted in the spread of HCV within the drug-user population. In fact, establishing more SSFs in Vancouver’s DTES would be a beneficial and fiscally responsible in addition to the publically funded health-care system. Based on the marginal counts, it should be noted that although expansion beyond the second SSF location may not provide the same economic return as the cumulative estimates, it may still be considered cost-effective given that the cumulative result was cost-effective beyond the seventh potential location.

Though not outlined in this analysis, there are several other benefits of opening a SSF that may add to the existing financial benefits of a SSF. One such benefit is the lowering of the risk of overdose, particularly for those smoking heroin and methamphetamine [63]. In British Columbia alone, 14 deaths have been attributed to heroin smoking [27]. Given the medical supervision of NIDUs, SSF has the potential to mitigate the risk of overdose deaths.

Another benefit of opening a SSF is the potential to increase detoxification and reduce risk behavior through education. Research indicates that NIDU will change their risk behavior when provided with appropriate education and treated with care [27]. Moreover, IDUs that regularly use the Vancouver’s Insite are more likely to initiate and maintain addiction treatment [4]. By visiting a SSF, people who use drugs may utilize various services such as mental health, counselling, and detoxification. Furthermore, SSFs can be expected to reduce public drug use in the same way that Vancouver’s Insite has been able to reduce public drug-use behavior of IDUs.

In summary, not only on the use of crack among drug users is on the rise but also the sharing of crack pipe has been increasing at an alarming rate in Vancouver. With recent research demonstrating the significant risk of disease transmission via oral smoking equipment, the current study determined whether the former unsanctioned SSF operated by the grassroot organization, VANDU, would cost less than the health-care consequences of not having such a program in Vancouver. The results indicated that the former facility not only saved taxpayers’ money but also deserved to be expanded instead of being forced to shut down. This information and analysis should be useful for policy makers who seek to find practical, cost-effective solutions to serious health-care problems in a climate of scarce public resources.

Abbreviations
VANDU: Vancouver Area Network of Drug Users; DTES: Downtown Eastside; SSF: supervised smoking facility; HCV: hepatitis C; IDU: injecting drug user; NIDU: non-injecting drug user; HIV: human immunodeficiency virus.

Competing interests
The authors declare that they have no competing interests.

Authors’ contributions
VANDU collected the data. EJ conducted the analysis and wrote the results, discussion, conclusion, and the references. Both authors (VANDU executive board) read and approved the final version of the paper.

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References
1. Werb D, DeBeck K, Kerr T, Li K, Montaner J, Wood E. Modelling crack cocaine use trends over 10 years in a Canadian setting. Drug Alcohol Rev 2010, 29:271–277.
2. Haydon E, Fischer B. Crack use as a public health problem in Canada call for an evaluation of safer crack use kits. Can J Public Health 2005, 96:185–188.
3. Fischer B, Manzoni P, Rehm J. Comparing injecting and non-injecting illicit opioid users in a multisite Canadian sample (OPICAN cohort). Eur Addict Res 2006, 12:230–239.
4. Persaud S, Tzemis D, Kuo M, Bungay V, Buxton JA. Controlling chaos: the perceptions of long-term crack cocaine users in Vancouver, British Columbia. Canada. J Addict 2013, 2013:1–9.
5. Office of Drugs and Alcohol Research and Surveillance Controlled Substances and Tobacco Directorate, Health Canada: Canadian Alcohol and Drug Use Monitoring Survey. Ottawa: Health Canada; 2010.
6. Ti L, Buxton J, Wood E, Zhang R, Montaner J, Kerr T: Difficulty accessing crack pipes and crack pipe sharing among people who use drugs in Vancouver. Canada. Subst Abuse Treat Prev Pol 2011, 6:34.

7. DeBeck K, Kerr T, Li K, Fischer B, Buxton J, Montaner J, Wood E: Smoking of cocaine as a risk factor for HIV infection among people who use injection drugs. Can J Infect Dis 2009, 18(1):585–589.

8. Fischer B, Rehm J, Patra J, Iakoubuk K, Haydon E, Tyndall M, El-Guebaly N: Crack across Canada: comparing crack users and crack non-users in a Canadian multi-city cohort of illicit opioid users. Addiction 2006, 101(12):1760–1770.

9. UNODC: World Drug Report 2007. Vienna: United Nations Office on Drugs and Crime, 2007.

10. Edlin BR, Irwin KL, Faruque S, McCoy CB, Word C, Serano Y, Inciardi JA, Bower BP, Schilling RF, Holmberg SD: Intersecting epidemics–crack cocaine use and HIV infection among inner-city young adults. N Engl J Med 1994, 331(21):1422–1427.

11. Buchanan D, Tooze JA, Shaw S, Kinzy M, Heimer R, Singer M: Demographic, HIV risk behavior, and health status characteristics of ‘crack’ cocaine injectors compared to other injection drug users in three New England cities. Drug Alcohol Depend 2006, 81(3):221–229.

12. Booth RE, Kwiatkowski CF, Chitwood DD: Substance use dynamics of life in the Downtown Eastside of Vancouver. J Acquir Immune Defic Syndr 2001, 28:273–278.

13. Cross JC, Johnson BD, Davis WR, Liberty H1: Supporting the habit: income generation activities of frequent crack users compared with frequent users of other hard drugs. Drug Alcohol Depend 2001, 64(2):191–201.

14. DeBeck K, Shannon K, Wood E, Li K, Montaner J, Kerr T: Income generating activities of people who inject drugs. Drug Alcohol Depend 2007, 91(1–50):56.

15. Booth RE, Kwiatkowski CF, Weissman G: Health-related service utilization and HIV risk behaviors among HIV infected injection drug users and crack smokers. Drug Alcohol Depend 1999, 55(1):219–78.

16. Malchyi L, Bungay V, Johnson J: Documenting practices and perceptions of ‘safer crack use’ a Canadian pilot study. Int J Drug Policy 2006, 17(5):571–572.

17. Tortu S, Neaigus A, McMahon J, Hagan D: Hepatitis C among noninjection drug users: a report. Subst Use Misuse 2001, 36:523–534.

18. Grymavith VA, Neaigus A, Miller M, Friedman SR, Des Jarlais DC: Risk correlates of prevalent HIV, hepatitis B virus, and hepatitis C virus infections among non-injection heroin users. J Acquir Immune Defic Syndr 2002, 30:448–456.

19. Hovey CJ, Fuller CM, Ompad DC, Galea S, Koblin B, Thomas D, Wahoo D: Association of sex, hygiene and drug equipment sharing with hepatitis C virus infection among non-injecting drug users in New York City. Drug Alcohol Depend 2005, 79:389–395.

20. Judd A, Hickman M, Rhodes T: Transmission of hepatitis C are noninjection cocaine users at risk? Subst Use Misuse 2002, 37:573–575.

21. Celentano D, Sherman SG: Commentary: the changing landscape of crack cocaine use and HIV infection. Can Med Assoc J 2009, 181(5):571–572.

22. Haydon E, Fischer B: Crack use as a public health problem in Canada: call for expanding Vancouver’s Insite. Urban Stud 2012, 49(8):3531–3544.

23. Jozaghi E, Reid AA, Andrews MA: A cost-benefit/cost-effectiveness analysis of proposed supervised injection facilities in Montreal, Canada. Subst Abus Treat Prev Policy 2013, 8:25.

24. Jozaghi E, Reid AA, Andrews MA, Janeua A: A cost-benefit/cost-effectiveness analysis of proposed supervised injection facilities in Ottawa, Canada. Subst Abus Treat Prev Policy 2014, 9:311–13.

25. Bayoumi AM, Zarcis GS: The cost-effectiveness of Vancouver’s supervised injection facility. Can Med Ass J 2008, 179(11):143–151.

26. Bayoumi AM, Strike C: Report of the Toronto and Ottawa Supervised Consumption Assessment Study. Toronto: University of Toronto; 2012.

27. Bravo MI, Royuela L, Fuente DL, Brugal MT, Barrio G, Salvany D: Use of supervised injection facilities and injection risk behaviours among young drug injectors. Addiction 2009, 104:614–619.

28. Ivins A, Roth E, Nakamura N, Kradjen M, Fischer B: Uptake, benefits of and barriers to safer crack use kit (SCUK) distribution programmes in Victoria, Canada—a qualitative exploration. Int J Drug Policy 2011, 22:292–300.

29. VANDU: Data Provided to the Researcher by the Vancouver Area Network of Drug Users. VANDU, Vancouver, 2013.

30. Mui M: Crackpipe distribution falls short. Four Hours Vancouver 2012, January 2. [http://vancouver24hrs.ca/News/local/2012/01/02/19191986.html]

31. Scheinmann R, Hagana H, Leulutu-Weinberger C, Stem R, Des Jarlais DC, Flora PL, Strauss S: Non-injection drug use and Hepatitis C Virus: a systematic review. Drug Alcohol Depend 2007, 89:1–12.

32. Grymavith VA, Neaigus A: The relationship of sexual dyad and personal network characteristics and individual attributes to unprotected sex among young injection drug users. AIDS Behav 2009, 13:196–206.

33. Fischer B, Rudzinski K, Ivins A, Gallupe O, Patra J, Kradjen M, Social, health and drug use characteristics of primary crack users in three mid-sized communities in British Columbia. Canada. Drug Educ Prev Pol 2010, 17(6):333–350.

34. Gilbert VL, Evans BG, Douglas S: HIV transmission among men who have sex with men through oral sex. Sex Transm Infect 2004, 80(3):24–28.

35. Kerr T, Wood E: What do you do when you hit rock bottom? Responding to drugs in the city of Vancouver. Int J Drug Pol 2006, 17(2):55–60.

36. Jozaghi E: An exploratory study involving the relationship between drug use and HIV risk behaviors among injection drug users. Int J Drug Policy 2004, 15(4):313–321.

37. Jozaghi E, Reid AA: A case study in transformation in the Downtown Eastside of Vancouver, Canada by peer injection drug users. Can J Crim J 2014, 56(5):535–594.

38. Culhan D: Their spirits live within us: aboriginal women in the Downtown Eastside Vancouver emerging into visibility. Am Ind Q 2003, 27(3):549–563.

39. The Safer Crack Use Coalition of Toronto: Crack, crack house sex, and HIV risk. Arch Sex Behav 1995, 24(3):249–269.

40. Kenny LE, Zarcis GS, Holodny M, Bendavid E, Owens DK, Brandeau ML: Cost-effectiveness of screening strategies for early identification of HIV and HCV infection in injection drug users. PLoS One 2012, 7(9):e45176.

41. John-Baptiste A, Yeung M, Leung V, van der Velde G, Krahn M: Cost-effectiveness of harm reduction interventions targeting substance users and other high-risk groups: a systematic review. Pharmacoeconomics 2012, 30(1):1015–1035.
58. Krajden M, Kuo M, Zagorski AM, Yu A, Krahn M: Health care costs associated with hepatitis C: a longitudinal cohort study. Can J Gastroenterol 2010, 24(12):717–726.

59. Jacobson IM, Brown RS Jr, Freilich B, Afdhal N, Kwo PY, Santoro J, Becker S, Wakil AE, Pound D, Godofsky E, Strauss R, Bernstein D, Flam A, Pauly MP, Mukhopadhyay P, Griffel LH, Braun CA, WIN-R Study Group: Peginterferon alfa-2b and weight-based or flat-dose ribavirin in chronic hepatitis C patients: a randomized trial. Hepatology 2007, 46(4):971–981.

60. Werb D, Wood E, Kerr T, Hershfield N, Palmer RW, Remis RS: Treatment costs of hepatitis C infection among injection drug users in Canada, 2006–2026. Int J Drug Policy 2011, 22:70–76.

61. Martin N, Vickerman P, Miners A, Foster GR, Hutchinson SJ, Goldberg DJ, Hickman M: Cost-effectiveness of hepatitis C virus antiviral treatment for injection drug user populations. Hepatology 2012, 55(1):49–57.

62. National Centre in HIV Epidemiology and Clinical Research: Epidemiological and Economic Impact of Potential Increased Hepatitis C Treatment Uptake in Australia. Sydney: The University of New South Wales; 2010.

63. Smith C: Baffling brain ailment hits heroin smokers, The Georgia Straight; 2004. http://www.straight.com/news/baffling-brain-ailment-hits-heroin-smokers.