Determinants of hospitalization for a cutaneous injection-related infection among injection drug users: a cohort study

Elisa Lloyd-Smith1,2, Evan Wood1,2,3, Ruth Zhang1,2, Mark W Tyndall1,2, Sam Sheps2, Julio SG Montaner1,2,3 and Thomas Kerr*1,2,3

Abstract
Background: Cutaneous injection-related infections (CIRI) are a primary reason individuals who inject drugs (IDU) are hospitalized. The objective of this study was to investigate determinants of hospitalization for a CIRI or related infectious complication among a cohort of supervised injection facility (SIF) users.

Methods: From 1 January 2004 until 31 January 2008, using Cox proportional hazard regression, we examined determinants of hospitalization for a CIRI or related infectious complication (based on ICD 10 codes) among 1083 IDU recruited from within the SIF. Length of stay in hospital and cost estimates, based on a fully-allocated costing model, was also evaluated.

Results: Among hospital admissions, 49% were due to a CIRI or related infectious complication. The incidence density for hospitalization for a CIRI or related infectious complication was 6.07 per 100 person-years (95% confidence intervals [CI]: 4.96 - 7.36). In the adjusted Cox proportional hazard model, being HIV positive (adjusted hazard ratio [AHR] = 1.79 [95% CI: 1.17 - 2.76]) and being referred to the hospital by a nurse at the SIF (AHR = 5.49 [95% CI: 3.48 - 8.67]) were associated with increased hospitalization. Length of stay in hospital was significantly shorter among participants referred to the hospital by a nurse at the SIF when compared to those who were not referred (4 days [interquartile range {IQR}: 2-7] versus 12 days [IQR: 5-33]) even after adjustment for confounders (p = 0.001).

Conclusions: A strong predictor of hospitalization for a CIRI or related infectious complication was being referred to the hospital by a nurse from the SIF. This finding indicates that nurses not only facilitate hospital utilization but may provide early intervention that prevents lengthy and expensive hospital visits for a CIRI or related infectious complication.

Background
Cutaneous injection-related infections (CIRI), which include cellulitis and abscesses, are among the primary causes of hospitalization among individuals who inject drugs (IDU) [1-4]. Hospitalizations due to CIRI carry considerable economic burden [3,5,6]. Complications of CIRI that are more likely to require hospitalization include, but are not limited to: osteomyelitis [7], bactere mia and sepsis [8,9], endocarditis [10,11], septic arthritis [7,12], ulcer [9], thrombophlebitis [13,14] and myositis [9].

A recent report by Hope et al. (2008) suggested that healthcare associated costs for CIRI among IDU in England were substantial, ranging from £15.5 million to £30.0 million per annum [5] ($27.1 to $52.4 million Canadian dollars). In 2001, Palepu et al. reported that of the IDU seen at St. Paul’s Hospital, an urban hospital in Vancouver, Canada, 35% had been hospitalized and a third of these hospitalizations were due to a CIRI or related infectious complication [1]. Hospitalization was expensive with hospital utilization cost per day reported to be $610 Canadian dollars (C$610) (95% confidence interval [CI]: C$576- C$645) [1].
North America’s first supervised injection facility (SIF) opened in Vancouver’s Downtown Eastside (DTES) in 2003. Within the SIF, IDU can inject pre-obtained drugs under the supervision of nurses. Individuals visiting the SIF are provided with sterile injecting equipment and emergency intervention in the event of an overdose, as well as primary medical care and addiction treatment, either on site or through referral. While several studies have pointed to the positive impact of the SIF on public disorder [15], HIV risk behaviour [16], management of overdose [17,18] and use of addiction treatment [19], its role with regard to hospitalization for CIRI remains unknown. However, recent research highlights that CIRI are a medical issue of concern among users of the SIF. Over a two year period, 6-10 per cent of IDU had a CIRI at time of interview [20]. In the present analysis, using longitudinal data, we examined predictors and cost of hospitalization for a CIRI or related infectious complication among IDU using the SIF.

Methods
Design and participants
The SIF in Vancouver is being evaluated through the Scientific Evaluation of Supervised Injection (SEOSI) cohort, which has been described in detail [21]. Briefly, the cohort was assembled through random recruitment of IDU from within the SIF. Random recruitment is based on inviting users of the SIF to be referred to the research study during random blocks of time. Among individuals who were recruited, a venous blood sample was drawn and an interviewer-administered questionnaire was conducted at baseline and at semi-annual follow-up visits. The informed consent agreement, obtained for all participants, included a request to link the SIF evaluation with administrative health databases. In Vancouver, hospitals are equipped with a database that tracks patient admission. The SIF is also equipped with a similar database. At the SIF, nursing care includes wound care, dressing changes, and measuring temperature. However, there are no physicians at the SIF. If microbiological investigation or medical treatment, including antibiotic prescription or intravenous antibiotic therapy is required, individuals at the SIF must be referred to the hospital or a medical clinic. In this study, a linkage of SEOSI participant data, SIF data and St. Paul’s Hospital inpatient data was performed. St. Paul’s Hospital is the major urban hospital serving the DTES community, one of Canada’s poorest postal codes. The University of British Columbia-Providence Health Care Research Ethics Board approved the present study.

Measurements
The start point for these analyses was enrollment into the SEOSI cohort and the endpoint was hospitalization for a CIRI or related infectious complication. The infectious complications included were based on previous literature [8,9]. The definition of the reason for hospitalization was based on International Classification of Diseases (ICD) 10 codes on patients’ hospital records and included: abscess (G061, G062, L020, L021, L022, L024, J851), cellulitis (L0300, L0310, L0311, L032, L0335, L038), osteomyelitis (M4620, M4625, M4629, M8617, M8618, M8661, M8663, M8666, M8681, M8691, M8695), staphylococcal infection 161 {(A490, A499, B956) including, sepsicaemia (A410, A412, A419) and Methicillin-Resistant Staphylococcus aureus (MRSA), (U000)}, endocarditis (I330), septic arthritis (M0000, M0002, M0004, M0005, M0006, M0008, M0009), ulcer (L089, L979), thrombophlebitis (I802, I808) and myositis (M6005, M6008). A few ICD-10 codes chosen suggest but do not require infectious etiologies (e.g., thrombophlebitis, myositis, and ulcer). The number of events refers to a CIRI or related infectious complication as a primary, secondary, tertiary, quaternary, or quinary diagnoses according to relevant ICD 10 codes unless otherwise specified.

We first examined the distribution and frequency of a CIRI or related infectious complication. Then, we evaluated length of stay in hospital among study participants and examined this outcome as a continuous variable in a linear model that adjusted for the following confounding variables: age, sex, HIV serostatus and SIF nurse referral. We then considered the cost of hospitalization, associated with CIRI, which was estimated at C$712 per hospital day, based on a fully-allocated costing model for the province of British Columbia from 2001 [1]. This estimate was updated to Canadian dollars in 2005 [22]. Fully-allocated costing includes costs associated with nursing care, medications, investigations, physician visits and length of stay as well as overhead, opportunity cost of hospital resources and a 5% depreciation of capital equipment [1]. Potential healthcare savings were estimated by multiplying the cost per day value (C$712) by the difference in number of days hospitalized among individuals with CIRI who were referred by a nurse within the SIF and those who were self referred to the hospital.

We investigated baseline characteristics stratified by hospitalization or not bivariately. Using Cox proportional hazard regression, we examined factors potentially associated with hospitalization. Variables considered for our analyses included: age; sex at birth (female vs. male); current residence in DTES (yes vs. no); living in unstable housing (yes vs. no); daily cocaine injection (yes vs. no); daily heroin injection (yes vs. no); daily speedball injection (yes vs. no); and HIV serostatus (positive vs. negative). As used previously, unstable housing was defined as living in a single room occupancy (SRO) hotel, shelter, recovery or transition house, jail, on the street, or having no fixed address [23]. Variables from the semi-annual...
questionnaire referred to behaviour that occurred in the last six months unless otherwise specified. We also examined whether a SIF nurse referral to the hospital was associated with hospitalization and, if so, whether length of stay was different given referral versus self referral. For this task, we conducted a record linkage matching their SEOSI identifying code to each participant’s record in the SIF database to determine nurse referral. Then, we linked his or her SEOSI identifying code with his or her unique personal health number to examine hospital records prior to the censor or event date.

Variable selection was based on previously published literature on a CIRI or related infectious complication and hospitalization among IDU [1,5,8,9,20]. Variables considered associated with hospitalization were analyzed in unadjusted analyses and adjusted Cox proportional hazard regression model. Time zero was defined as the date of recruitment into the SEOSI study for all participants and participants not hospitalized at St. Paul’s Hospital were censored as of 31 January 2008. All behavioral variables were treated as time-updated covariates based on semi-annual follow-up data. The multivariate model was fit using a fixed model whereby we included all variables that were statistically significant at the \( p < 0.05 \) level in univariate analyses. All statistical analyses were performed using SAS 8.0 (Cary, NC) and all \( p \)-values were two-sided.

**Results**

During the study period (1 January 2004 to 31 January 2008), 1083 individuals were recruited into the SEOSI cohort and 901 (83%) reported at least one follow-up visit. The median age among SEOSI participants was 38.4 years (interquartile range [IQR]: 32.7-44.3) and 314 (29%) were female. The median follow-up duration after recruitment into the cohort was 21.4 months (IQR: 13.1-24.6). Among the study period, 99 (9%) participants were admitted to St. Paul’s Hospital, yielding an incidence density of hospitalization for a CIRI or related infectious complication of 6.07 per 100 person-years (95% CI: 4.96 - 7.36). Among all hospital admissions among SEOSI participants, 49% (216 of 442) were related to a CIRI or related infectious complication. A total of 216 hospitalization events occurred among 99 individuals for a CIRI or related infectious complication. Fifteen persons had missing data on HIV serostatus and were excluded from analyses. Therefore, all results on HIV were based on a sample size of 1068. Participants who had been referred to the hospital by an SIF nurse had a significantly shorter length of stay in hospital as compared to those who were not referred (4 days [IQR: 2-7], 12 days [IQR: 5-33]). The eight day reduction in the length of hospital stay remained significant (\( p = 0.001 \)) after adjustment for confounding variables. Considering the total potential healthcare saving based on a fully allocated hospital cost per day calculation, each referral from the SIF would have resulted in a saving of C$5,696 (IQR: C$2, 136 - C$18, 512).

The factors associated with an increased risk of hospitalization after recruitment into the SEOSI cohort are shown in Table 4. In the multivariate model being HIV positive (adjusted hazard ratio [AHR] = 1.79 [95% CI: 1.16 - 2.75]) and being referred to the hospital by a nurse at the SIF (AHR = 5.38 [95% CI: 3.39 - 8.55]) were positively and independently associated with an increased likelihood of hospitalization.

| Classification† | First event \( n \) (%) | Total events \( n \) (%) |
|-----------------|-----------------|-----------------|
| Cellularis      | 33 (33)         | 59 (27)         |
| Abscess         | 14 (14)         | 26 (12)         |
| Osteomyelitis   | 10 (10)         | 39 (18)         |
| Staphylococcal infection | 22 (22) | 42 (17)         |
| Endocarditis    | 9 (9)           | 24 (12)         |
| Septic arthritis | 7 (7)        | 12 (6)          |
| Ulcer           | 1 (1)           | 6 (3)           |
| Thrombophlebitis | 1 (1)         | 4 (2)           |
| Myositis        | 2 (2)           | 4 (2)           |

† Classification is based on International Classification of Diseases 10 codes
Discussion
In the present study, being referred by a nurse at a SIF was independently associated with an elevated rate of hospitalization. Importantly, participants who had been referred to the hospital by a nurse at the SIF had a significantly shorter length of stay in hospital despite adjustment for HIV infection and other potential confounders. This finding indicates that nurses facilitate hospital utili-

| Table 2: Distribution of hospitalizations for a cutaneous injection-related infection or related infectious complication*.

| Primary diagnosis | +1 | +2 | +3 | +4 | +5 | +6 | +7 | +8 | +9 | +1&2 | +1&4 |
|------------------|----|----|----|----|----|----|----|----|----|------|------|
| 1cellulitis       | 45 | -  | 3  | 0  | 5  | 0  | 0  | 2  | 1  | 0    | -    |
| 2abscess         | 9  | 1  | -  | 0  | 5  | 0  | 0  | 0  | 0  | -    | 3    |
| 3osteomyelitis   | 14 | 0  | 5  | 1  | 0  | 0  | 0  | 1  | 2  | 0    | 1    |
| 4staph.infection | 18 | 3  | 0  | 1  | 4  | 0  | 0  | 0  | 1  | -    | -    |
| 5endocarditis    | 12 | 0  | 1  | 0  | 5  | -  | 0  | 0  | 0  | 1    | -    |
| 6septic arthritis| 8  | 0  | 2  | 0  | 1  | 0  | -  | 0  | 0  | 0    | 1    |
| 7ulcer           | 3  | 0  | 0  | 0  | 0  | 0  | 0  | 0  | 0  | -    | 0    |
| 8thrombophlebitis| 3  | 0  | 0  | 1  | 0  | 0  | 0  | 0  | 0  | -    | 0    |
| 9myositis        | 1  | 0  | 0  | 0  | 3  | 0  | 0  | 0  | -  | 0    | 0    |

+1&7  +2&4  +3&4  +3&7  +4&7  +5&4  +9&4  + more than two diseases

| Primary diagnosis | +1&7 | +2&4 | +3&4 | +3&7 | +4&7 | +5&4 | +9&4 | + more than two diseases |
|------------------|------|------|------|------|------|------|------|--------------------------|
| 1cellulitis       | -    | 0    | 0    | 1    | 2    | 0    | 0    | 0                        |
| 2abscess         | 0    | -    | 2    | 0    | 0    | 1    | 1    | 4                        |
| 3osteomyelitis   | 1    | 3    | -    | -    | 2    | 0    | 0    | 5                        |
| 4staph.infection | 0    | -    | -    | 0    | -    | -    | -    | 15                       |
| 5endocarditis    | 0    | 0    | 0    | 0    | 0    | -    | 0    | 5                        |
| 6septic arthritis| 0    | 0    | 0    | 0    | 0    | 0    | 0    | 0                        |
| 7ulcer           | -    | 0    | 0    | -    | 0    | 0    | 0    | 2                        |
| 8thrombophlebitis| 0    | 0    | 0    | 0    | 0    | 0    | 0    | 0                        |
| 9myositis        | 0    | 0    | 0    | 0    | 0    | -    | 0    | 0                        |

*According to ICD 10 codes. ICD 10 codes categorized into 9 diseases and listed from 1 to 9. First column refers to when disease listed as primary diagnosis. Other columns refer to multiple diseases reported, irrespective of whether event was primary, secondary, tertiary, quaternary, or quinary.

Table 3: Frequency of hospitalizations for a cutaneous injection-related infection or related infectious complication*.

|                  | Once | Twice | Three times | Four times | Five plus times | Total |
|------------------|------|-------|-------------|------------|----------------|-------|
| 1cellulitis      | 19   | 7     | 3           | 0          | 2              | 59    |
| 2abscess         | 10   | 5     | 2           | 0          | 0              | 26    |
| 3osteomyelitis   | 5    | 2     | 2           | 0          | 3              | 39    |
| 4staph.infection | 10   | 2     | 0           | 3          | 2              | 42    |
| 5endocarditis    | 11   | 0     | 1           | 0          | 1              | 24    |
| 6septic arthritis| 6    | 3     | 0           | 0          | 0              | 12    |
| 7ulcer           | 1    | 0     | 1           | 0          | 1              | 6     |
| 8thrombophlebitis| 1    | 0     | 1           | 0          | 0              | 4     |
| 9myositis        | 3    | 0     | 1           | 0          | 0              | 4     |

*According to ICD 10 codes. Diseases listed from 1 to 9. ICD 10 codes categorized into 9 diseases and listed from 1 to 9. Data relates to number of times an individual developed a disease, irrespective of whether event was coded as primary, secondary, tertiary, quaternary, or quinary.
zation as well as providing early intervention that prevents lengthy and expensive hospital visits for a CIRI or related infectious complication. According to the cost savings calculation based on the length of stay reduction, SIF nurse referral resulted in a minimum saving of C$2,136 per admission. However, it is important to note that variability in treatment costs exists in the CIRI and related infectious complications included in this investigation.

Using the same cohort, the risk factors for the outcome of developing a CIRI have been reported to include being female, living in unstable housing, borrowing syringes, requiring help injecting and injecting cocaine daily [20]. Some factors differ from risk factors in other cities. For example, research from San Francisco reports an associated between developing a CIRI and frequent black tar heroin injection [2] whereas in Vancouver, frequent cocaine injection has been associated with developing a CIRI [20]. The present study evaluated the predictors of the outcome of hospitalization for a CIRI or related infectious complication. The predictors of hospitalization for a CIRI or related infectious complication included being HIV positive and referred to the hospital by a nurse at the SIF. Future research investigating outcomes, including mortality from a CIRI or related infectious complication among those referred compared to those not referred to the hospital by a nurse at the supervised injection facility would support our understanding of the potential impact of nurse referral. Interestingly, being HIV positive was not a risk factor for the development of CIRI but was a predictor of hospitalization for a CIRI or related infectious complication.

As noted above, an increased risk of hospitalization for a CIRI or related infectious complication was observed among HIV positive participants. This finding is consistent with previous research on hospitalization among IDU in this setting [1]. Individuals with HIV may be more likely to be treated as an inpatient as opposed to an outpatient (i.e., HIV positive patients with abscesses may be more likely to get admitted to the hospital than HIV negative patients with abscesses). A potentially elevated susceptibility to bacterial infections [8,24] as well as known high-risk drug injection practices in this subpopulation [25,26] may related to why HIV positive participants in this study had elevated rates of hospitalization. Further elucidation of this finding with a larger sample of HIV positive IDU is necessary to examine this important research question.

After controlling for factors that are known to be associated with hospitalization [1,27], referral from an SIF nurse remained a strong predictor of hospitalization. Findings from this study confirm existing qualitative research indicating that nurses at the SIF provide CIRI-related care and play a key role in enhancing access to medical care among participants who have infections or injuries of greater severity [28]. Hospital care for more severe forms of CIRI is essential since local hospitals provide treatment to CIRI not available at the SIF (e.g., incision and drainage of an abscess, intravenous antibiotic therapy as well as diagnostic tools and therapy appropriate for complications). Further, given that the average length of stay in hospital for those referred to the hospital by a nurse at the SIF was significantly shorter, it may be that nurses at the SIF are helping reduce the incidence of late presentation of complications, such as osteomyelitis, that are often lengthy and are particularly costly to the healthcare system [7]. Although our study design does not allow us to infer causation, it is noteworthy that this association persisted in multivariate analyses. This may reflect prompt referral as well as the preventative effect of the SIF on serious infections [16].

There are several examples of efficient and cost effective community-based treatment services that may inform changes in the provision of care that is required in order to reduce the incidence of hospitalization of a CIRI or related infectious complication in our setting. In its first year of operation, the Integrated Soft Tissue Infection Services (ISIS) Clinic in San Francisco, California, resulted in a 47% decrease in surgical service admissions and an estimated savings of over $8.0 (C$8.4) million for costs related to CIRI [29]. In addition, a wound management clinic operated in conjunction with a syringe exchange program in Oakland, California [30]. It was found that the average cost per individual treated at this wound management clinic was $5.0 (C$5.2), substantially lower than equivalent hospital costs of $185.0 (C$193.4) and $360.0 (C$376.3) [30]. Given the success other cities have had in implementing additional medical treatment for CIRI in the community, we recommend initiating provision of on-site incision and drainage for abscesses and administration of antibiotic therapy at the SIF or in a nearby community setting. Further, to prevent development of CIRI, it may be of value to screen and treat for MRSA at the SIF or at the SEOSI research site. In our setting, expanding the capacity to provide primary care in an integrated manner is warranted.

There are limitations of the present study to consider when interpreting this data. Firstly, St. Paul’s Hospital was the only facility linkage that was conducted. However, this is the primary hospital serving the SIF catchment area and is accessed extensively by the IDU population in the DTES [1,27,31]. Secondly, our study relies on self report to obtain drug use and other behavioural variables. However, self report among IDU is considered valid [32] and hospital utilization and nurse referral were accessed directly from databases within the hospital and at the SIF, respectively. Thirdly, a limitation of this analysis is that a low sample size of HIV positive participants precluded
our ability to conduct further analyses on this association. Additional research is required with a larger sample of HIV positive individuals to better elucidate this finding. Fourthly, additional research is required to better understand why a shorter length of stay is seen among SEOSI participants referred by a nurse at the SIF to the hospital. It is possible that nurses may be referring to the hospital SEOSI participants that require shorter lengths of stay (e.g., abscess versus osteomyelitis). However, in our study, individuals with an ED visit who were referred by a nurse were significantly more likely to be hospitalized within three days. Similarly, Binswanger et al. found ED use for CIRI to be associated with hospitalization and also death [33].

**Conclusion**

In summary, we found high levels of hospitalization for a CIRI or related infectious complication among local IDU. Being HIV positive and being referred to the hospital by a nurse at the SIF, as opposed to self referral to the hospital, were both independently and positively associated with an increased likelihood of hospitalization. Participants who had been referred to the hospital by a nurse at the SIF had a significantly shorter length of stay in hospital as compared to those who were not. These findings indicate that nurses at the SIF play a critical role in terms of referring individuals who require hospitalization for a CIRI or related infectious complication to the hospital, which may result in shorter and less expensive hospital visits. Expanded management of CIRI in the community may reduce the need for referral to the hospital, further reducing the cost of caring for this common clinical problem among IDU.

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

**Authors’ contributions**
ELS conceived and designed the study and drafted the manuscript. RZ and ELS performed the statistical analyses. ELS, TK, EW, MT contributed to the design and coordination of the study. All authors provided assistance with interpretation of the results to the drafts of the manuscript and read and approved the final manuscript.

**Acknowledgements**
The authors wish to thank the participants in SEOSI and the staff at Insite, the Portland Hotel Society, and Vancouver Coastal Health (Chris Buchner, David Marsh and Heather Hay). We also thank all current and past SEOSI staff. We would also like to thank Deborah Graham, Leslie Rae, Caitlin Johnston, Steven Kain and Calvin Lai for their research assistance. Viviane Dias Lima and Karissa Johnston contributed statistical expertise. The evaluation of the SIF was originally made possible through a financial contribution from Health Canada, although the views expressed here do not reflect the official policies of Health Canada. The evaluation is currently supported by the Canadian Institutes of Health Research (grants HPR-85526 and RAA-79918) and Vancouver Coastal Health. TK, MT and ELS are supported by the Michael Smith Foundation for Health Research; TK and ELS are supported by the Canadian Institutes of Health Research.

| Variable                  | Unadjusted Hazard Ratio (HR) | Unadjusted p-value | Adjusted Hazard Ratio (AHR) | Adjusted p-value |
|---------------------------|------------------------------|--------------------|-----------------------------|------------------|
| Age                       | 0.98 (0.96 - 1.01)           | 0.146              | 1.36 (0.90 - 2.05)           | 0.139            |
| Sex                       |                               |                    |                             |                  |
| (Female vs Male)          | 1.59 (1.07 - 2.39)           | 0.024              | 1.26 (0.79 - 2.02)           | 0.328            |
| Unstable housing*         | 1.65 (1.08 - 2.53)           | 0.021              | 1.26 (0.79 - 2.02)           | 0.328            |
| Cocaine injection*        | 1.75 (1.17 - 2.62)           | 0.006              | 1.26 (0.79 - 2.02)           | 0.328            |
| Speedball injection*      | 1.90 (1.15 - 3.14)           | 0.012              | 1.19 (0.69 - 2.07)           | 0.528            |
| HIV serostatus            |                               |                    |                             |                  |
| (Positive vs Negative)    | 2.12 (1.39 - 3.24)           | <0.001             | 1.79 (1.16 - 2.75)           | 0.008            |
| Hospital referral†        | 2.41 (1.55 - 3.77)           | <0.001             | 5.38 (3.39 - 8.55)           | <0.001           |

*Behaviour refers to activities in the last 6 months. †Indicates data derived from SIF database and by a study nurse. CI = confidence interval.
References

1. Palepu A, Tyndall MW, Leon H, Muller J, O'Shaughnessy MV, Schechter MT, et al.: Hospital utilization and costs in a cohort of injection drug users. CMAJ 2001, 165:415-420.

2. Murphy E, DeVita D, Liu H, Vittinghoff E, Leung P, Ciccarone DH, et al.: Frequency, factors and costs associated with injection site infections: findings from a national multi-site survey of injecting drug users in England. BMC Infect Dis 2008, 8:

3. Ciccarone D, Bamberger JD, Kral AH, Edlin BR, Hobart CJ, Moon A, et al.: Soft tissue infections among injection drug users - San Francisco, California, 1996-2000. MMWR Morb Mortal Wkly Rep 2001, 50:381-384.

4. Takahashi TA, Baenentstein A, Binswanger I, Bradley K, Merrill JO. Predictors of hospitalization for injection drug users seeking care for soft tissue infections. J Gen Intern Med 2007, 22:382-388.

5. Hope V, Kimber J, Vickerman P, Hickman M, Ncube F. Methadone maintenance treatment and attendance at attendance at a medically supervised injecting facility: a qualitative study. Drug Alcohol Depend 2001, 62-66.

6. DiNubile MJ, Lipsky BA. Complicated infections of the skin and skin structures: when the infection is more than skin deep. J Antimicrob Chemother 2004, 53:i37-i40.

7. Kak V, Chandrasekar PH: Methanobrevibacter smithii is a normal inhabitant of the oral cavity. J Clin Microbiol 2007, 45:2189-2191.

8. Schulz S, Beckenbach C, Philipp M, Hengstmann J. Color coded duplex sonography of inguinal vessels in i.v. drug addicts. Radiographics 2003, 23:721-728.

9. Klotz M, Krausshofer H. Epidemiology of endocarditis caused by Staphylococcus aureus. J Antimicrob Chemother 2002, 50:535-538.

10. Hope V, Kimber J, Vickerman P, Hickman M, Ncube F. Methadone maintenance treatment and attendance at attendance at a medically supervised injecting facility: a qualitative study. Drug Alcohol Depend 2001, 62-66.

11. Ngaage DL, Cowen ME. Right ventricular needle embolus in an injecting drug user: the need for early removal. Emerg Med J 2001, 18:500-501.

12. Ebright JR, Pieter B. Skin and soft tissue infections in injection drug users. Infect Dis Clin North Am 2002, 16:681-695.

13. Schulz S, Beckenbach C, Philipp M, Hengstmann J. Color coded duplex sonography of inguinal vessels in i.v. drug addicts. Radiographics 2003, 23:721-728.

14. Klotz M, Krausshofer H. Epidemiology of endocarditis caused by Staphylococcus aureus. J Antimicrob Chemother 2002, 50:535-538.

15. de Vries RS, Nullens AC, Remon JA, van Loon T. Complicated infections of the skin and skin structures: when the infection is more than skin deep. J Antimicrob Chemother 2004, 53:i37-i40.

16. Kak V, Chandrasekar PH: Methanobrevibacter smithii is a normal inhabitant of the oral cavity. J Clin Microbiol 2007, 45:2189-2191.

17. Schulz S, Beckenbach C, Philipp M, Hengstmann J. Color coded duplex sonography of inguinal vessels in i.v. drug addicts. Radiographics 2003, 23:721-728.

18. Klotz M, Krausshofer H. Epidemiology of endocarditis caused by Staphylococcus aureus. J Antimicrob Chemother 2002, 50:535-538.

19. de Vries RS, Nullens AC, Remon JA, van Loon T. Complicated infections of the skin and skin structures: when the infection is more than skin deep. J Antimicrob Chemother 2004, 53:i37-i40.

20. Kak V, Chandrasekar PH: Methanobrevibacter smithii is a normal inhabitant of the oral cavity. J Clin Microbiol 2007, 45:2189-2191.

21. Schulz S, Beckenbach C, Philipp M, Hengstmann J. Color coded duplex sonography of inguinal vessels in i.v. drug addicts. Radiographics 2003, 23:721-728.

22. Klotz M, Krausshofer H. Epidemiology of endocarditis caused by Staphylococcus aureus. J Antimicrob Chemother 2002, 50:535-538.

23. de Vries RS, Nullens AC, Remon JA, van Loon T. Complicated infections of the skin and skin structures: when the infection is more than skin deep. J Antimicrob Chemother 2004, 53:i37-i40.

24. Kak V, Chandrasekar PH: Methanobrevibacter smithii is a normal inhabitant of the oral cavity. J Clin Microbiol 2007, 45:2189-2191.

25. Schulz S, Beckenbach C, Philipp M, Hengstmann J. Color coded duplex sonography of inguinal vessels in i.v. drug addicts. Radiographics 2003, 23:721-728.

26. Klotz M, Krausshofer H. Epidemiology of endocarditis caused by Staphylococcus aureus. J Antimicrob Chemother 2002, 50:535-538.

27. de Vries RS, Nullens AC, Remon JA, van Loon T. Complicated infections of the skin and skin structures: when the infection is more than skin deep. J Antimicrob Chemother 2004, 53:i37-i40.

28. Kak V, Chandrasekar PH: Methanobrevibacter smithii is a normal inhabitant of the oral cavity. J Clin Microbiol 2007, 45:2189-2191.

29. Schulz S, Beckenbach C, Philipp M, Hengstmann J. Color coded duplex sonography of inguinal vessels in i.v. drug addicts. Radiographics 2003, 23:721-728.

30. Klotz M, Krausshofer H. Epidemiology of endocarditis caused by Staphylococcus aureus. J Antimicrob Chemother 2002, 50:535-538.

31. de Vries RS, Nullens AC, Remon JA, van Loon T. Complicated infections of the skin and skin structures: when the infection is more than skin deep. J Antimicrob Chemother 2004, 53:i37-i40.

32. Kak V, Chandrasekar PH: Methanobrevibacter smithii is a normal inhabitant of the oral cavity. J Clin Microbiol 2007, 45:2189-2191.

33. Schulz S, Beckenbach C, Philipp M, Hengstmann J. Color coded duplex sonography of inguinal vessels in i.v. drug addicts. Radiographics 2003, 23:721-728.

34. Klotz M, Krausshofer H. Epidemiology of endocarditis caused by Staphylococcus aureus. J Antimicrob Chemother 2002, 50:535-538.

35. de Vries RS, Nullens AC, Remon JA, van Loon T. Complicated infections of the skin and skin structures: when the infection is more than skin deep. J Antimicrob Chemother 2004, 53:i37-i40.

36. Kak V, Chandrasekar PH: Methanobrevibacter smithii is a normal inhabitant of the oral cavity. J Clin Microbiol 2007, 45:2189-2191.

37. Schulz S, Beckenbach C, Philipp M, Hengstmann J. Color coded duplex sonography of inguinal vessels in i.v. drug addicts. Radiographics 2003, 23:721-728.

38. Klotz M, Krausshofer H. Epidemiology of endocarditis caused by Staphylococcus aureus. J Antimicrob Chemother 2002, 50:535-538.

39. de Vries RS, Nullens AC, Remon JA, van Loon T. Complicated infections of the skin and skin structures: when the infection is more than skin deep. J Antimicrob Chemother 2004, 53:i37-i40.

40. Kak V, Chandrasekar PH: Methanobrevibacter smithii is a normal inhabitant of the oral cavity. J Clin Microbiol 2007, 45:2189-2191.