Injecting drug use in Brighton, Liverpool, and London: best estimates of prevalence and coverage of public health indicators

Matthew Hickman, Vanessa Higgins, Vivian Hope, Mark Bellis, Kate Tilling, Angeline Walker, John Henry

Study objective: To estimate the prevalence of injecting drug use (IDU) in three cities in England and to measure the coverage of key public health indicators.

Design: Capture-recapture techniques with covariate effects.

Setting: Liverpool, Brighton, and 12 London boroughs, 2000/01.

Participants: IDU collated and matched across five data sources—community recruited survey, specialist drug treatment, arrest referral, syringe exchange, and accident and emergency—896 in Brighton, 1224 in Liverpool, and 6111 in London.

Main results: It is estimated that in 2000/01 the number and prevalence of IDU aged 15–44 was 2300 (95%CI 1500 to 3700) and 2.0% (95%CI 1.3% to 3.2%) in Brighton; 2900 (95%CI 2500 to 5000) and 1.5% (95%CI 1.3% to 2.6%) in Liverpool; 16 700 (95%CI 13 800 to 21 600) and 1.2% (95%CI 1.0% to 1.6%) in 12 London boroughs; with a prevalence of 1.7% (95%CI 1.2% to 3.3%) in inner London. It is estimated that: less than one in four IDU are in treatment in the three areas; syringe exchange programmes covered about 25% of injections in Brighton and Liverpool and 20% in London; and that the annual opioid mortality rate among IDU was 2% in Brighton compared with less than 1% in Liverpool and London.

Conclusions: Credible estimates of the prevalence of injecting drug use (and key public health indicators) can be determined using covariate capture-recapture techniques. These suggest that: targets to double the number in treatment are possible: syringe distribution should be increased; and further attention, especially in Brighton, given to reducing overdose mortality.

Knowledge of the prevalence of injecting drug use is critical to monitoring the United Kingdom’s current drug strategy that aims to increase the proportion of problem drug users in treatment; and UNAIDS targets that aim to measure the proportion of injecting drug users in contact with prevention services. In the UK, arguments over the extent of problem drug use, the coverage of treatment, and by extension whether “we have lost the drugs war” have also recently become front page news. But the evidence to support or counter these accusations is not available.

In the UK as in many other countries routine statistics on injecting or other problematic drug use do not regularly provide estimates of prevalence. Neither can “direct” estimates through population surveys provide reliable data because of multiple response biases. For instance, the 2001 British Crime Survey, which has a sample size of almost 33 000, found less than 50 people reporting that they used heroin in the past month, and the resulting population estimates (33 000, range 19 000 to 53 000) were implausible, falling short of the numbers of heroin users presenting to GPs and drug treatment agencies.

Indirect methods offer an alternative approach, developed for sampling rare or elusive populations (animal and human) in situations where there is no sampling frame and direct methods are impracticable. Capture-recapture is one such method: developed by animal ecologists, adopted first by demographers, and then used and promoted by epidemiologists. Capture-recapture has adjusted for the inevitable under-ascertainment of surveys and surveillance data for a variety of conditions including diabetes, congenital disorders, infectious disease, and injury. Its use for estimating the prevalence of “addiction” was first recognised by Bishop et al, and one off estimates have been generated for many cities worldwide.

In this study we focus on three cities in England (Brighton, Liverpool, and part of London), using new developments in the analysis of capture-recapture studies to estimate the number of injecting drug users (IDUs).

METHODS

Overview

In capture-recapture, two, but ideally three, or more data sources that identify the target population (that is, IDUs) are collected and matched using name, or often, as in our case, a set of “attributors” (person initials, date of birth, and sex) to identify the number of individuals in the whole dataset and the number of matches (that is, the number of people that occur in more than one data source). An equivalence is assumed between the level of overlap between the data sources (that is, the proportion of subjects in two or more data sources) and the sampling intensity (that is, the proportion of the total population recorded by the data sources). Simple equations or statistical models fitted to the observed data are used to estimate the number of injecting drug users who appear in none of the data sources, which combined with the observed number generates the prevalence estimate. The data sources and the statistical techniques used are summarised below.

Abbreviations: IDU, injecting drug use; SEP, syringe exchange programme; A&E, accident and emergency
The study was granted ethical approval by the Scottish Multi-Centre Research Ethics Committee, and 12 local research ethics committees.

Data sources
The target population was injecting drug users aged 15–44 resident in Liverpool city (population 195 000 aged 15–44), Brighton and Hove (117 000 aged 15–44), and 12 London boroughs comprising about one third of London and 60% of inner London (see footnote to table 1, population 1.36 million aged 15–44) during the period April 2000 to end March 2001. Data including matching attributors’ drug, and area of residence were collected from five sources:

1. Arrest referral: routine monitoring data were available electronically on assessments by arrest referral workers (who are located in police custody suites throughout England to assess drug problems of arrestees and refer, if appropriate, to a specialist drug agency).

2. Drug treatment: electronic data were retrieved from local databases and the National Drug Treatment Monitoring System, which collect reports of new and ongoing clients at specialist drug treatment agencies and a few primary care services.


4. Accident and emergency (A&E): attendees because of an overdose were identified through A&E computers and data extracted from patient notes from one hospital each in Liverpool and Brighton and 10 in London.

5. Community recruited survey: injecting drug users resident in the study area were interviewed using indigenous fieldworkers recruiting subjects from the street or social networks: 96 in Brighton, 151 in Liverpool, and 436 in London.

The A&E and community survey data sources in Brighton and Liverpool were combined because they were small, and in London they were combined further with arrest referral in order to cover the geographical area. Criteria for matches (subjects on more than one dataset) were: identical initials, date of birth, and gender; initials reversed, identical date of birth, and gender; number substitution in date of birth, identical initials, and gender. Three public health indicators were collected from a variety of sources (see footnote to table 3): number of specialist drug treatment; annual number of syringes distributed through syringe exchange services; and opioid overdose deaths.

Statistical techniques
The assumptions underpinning capture-recapture and the estimation equations are described in detail elsewhere. Two key biases that need to be controlled are “dependency” and “heterogeneity”. Dependencies arise if a person in one data source is more or less likely to be on another data source (for example, IDU in specialist drug treatment are more likely to be registered with a local syringe exchange programme than IDU not in treatment), and are simply tested by fitting an interaction between the data sources with the Poisson model and calculating whether this improves the fit of the model. With four data sources there are 114 potential models from “independence” (no interactions) to all three way interactions between the data sources.

Heterogeneity arises if the probability of being captured by a source is not equal for all members of the target population. Most health and criminal justice data will exhibit heterogeneity. For example, there were proportionally more males and subjects were younger in the arrest referral data source compared with the other data sources. The traditional capture-recapture approach has been to stratify the dataset to form homogeneous subsets, and then to carry out capture-recapture estimations separately on each subset. However, this increases the number of models (for each age group and gender, etc) used to provide estimates, involving multiple significance testing, and in small subsets (for example, young women) reducing the statistical power to distinguish the best model. Covariate capture-recapture techniques developed by Tilling et al adjust for heterogeneity more efficiently within fewer models through expanding the observed dataset to include covariates. Covariates (age group and sex) were both included as categorical values, given no reason to assume that the probability of capture varied linearly with age. In effect, by including all interactions between covariates the estimates will equate to a stratified solution, which also assumes no high order interactions, but it tests whether a different stratified model is justified and so leads to a more parsimonious model.

In this study, covariates (age group and gender) were created and included in the model by generating multiple dummy variables denoting dependencies between data sources and between covariates and the data sources, and Poisson models were fitted to the data sources and covariates and the overlap between them. Covariate analysis followed stratified analysis where within each strata the best fitting model, with the fewest parameters and smallest difference between observed and predicted values, was selected on the basis of standard information criterion. In the stratified analysis these included the likelihood ratio test (LRT) for competing models with different numbers of interactions, and the Bayes information criterion (also known as Drapers information criterion, DIC) or Akaike information criterion (AIC) (data not shown). We adopted this approach to select the simplest model given a variety of information criterion, though some analyses in the literature recommend the AIC as preferable. The covariate analysis began with all the interactions identified by the stratified analysis, and then selected the most parsimonious model (that is, with fewest parameters) based on the lowest AIC. The “best fitting poisson model” was used to generate estimates of the unobserved number of male and female IDU aged 15–29 and 30–44. Ninety five per cent confidence intervals (CI) were generated using bootstrap methods. These deal with model uncertainty, but assume that the model is correct, which we preferred to a solution that weights different models.

Policy implications

- Evidence on the number of injecting drug users can be obtained, but these studies are labour intensive. Introducing “prevalence estimation” as an objective of routine surveillance and condition of government investment is the answer to how the situation can be improved.

- Better quality data are required on the number of people in treatment, to make best use of evidence on the prevalence of injecting drug use.

- Syringe distribution at least in three cities in England may not be adequate to prevent transmission of hepatitis C.

- Identifying effective interventions to prevent and/or reverse transitions to injection drug use must become a critical public health action.
Table 1

<table>
<thead>
<tr>
<th>Data sources</th>
<th>Brighton</th>
<th>Liverpool</th>
<th>London</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Drug treatment</strong></td>
<td><strong>Syringe</strong></td>
<td><strong>Exchange programme</strong></td>
<td><strong>Survey</strong></td>
</tr>
<tr>
<td><strong>Male</strong></td>
<td><strong>Male</strong></td>
<td><strong>Male</strong></td>
<td><strong>Male</strong></td>
</tr>
<tr>
<td><strong>Females</strong></td>
<td><strong>Females</strong></td>
<td><strong>Females</strong></td>
<td><strong>Females</strong></td>
</tr>
<tr>
<td><strong>Total</strong></td>
<td><strong>Total</strong></td>
<td><strong>Total</strong></td>
<td><strong>Total</strong></td>
</tr>
<tr>
<td><strong>15–29</strong></td>
<td><strong>15–29</strong></td>
<td><strong>15–29</strong></td>
<td><strong>15–29</strong></td>
</tr>
<tr>
<td><strong>30–44</strong></td>
<td><strong>30–44</strong></td>
<td><strong>30–44</strong></td>
<td><strong>30–44</strong></td>
</tr>
<tr>
<td><strong>Yes</strong></td>
<td><strong>Yes</strong></td>
<td><strong>Yes</strong></td>
<td><strong>Yes</strong></td>
</tr>
<tr>
<td><strong>No</strong></td>
<td><strong>No</strong></td>
<td><strong>No</strong></td>
<td><strong>No</strong></td>
</tr>
<tr>
<td><strong>Yes</strong></td>
<td><strong>Yes</strong></td>
<td><strong>Yes</strong></td>
<td><strong>Yes</strong></td>
</tr>
<tr>
<td><strong>No</strong></td>
<td><strong>No</strong></td>
<td><strong>No</strong></td>
<td><strong>No</strong></td>
</tr>
</tbody>
</table>

The estimated prevalence varied by age and gender in all three cities. In Brighton, the estimated prevalence was 2.4% among men aged 15 to 29, 3.2% among women aged 15 to 29, and 3.2% among men aged 30 to 44. In Liverpool, the estimated prevalence was 2.4% among men aged 15 to 29 and 2.8% among women aged 15 to 29. In London, the estimated prevalence was 1.5% among men aged 15 to 29 and 1.5% among women aged 15 to 29.

The London estimate, when divided into inner and outer London, was 1.2% (95% CI 1.0% to 1.5%). The estimate for inner London was 1.2% (95% CI 1.0% to 1.5%) and the estimate for outer London was 1.2% (95% CI 1.0% to 1.5%).

The estimated prevalence of injecting drug users varied by age and gender in all three cities. In Brighton, the estimated prevalence was 2.4% among men aged 15 to 29, 3.2% among women aged 15 to 29, and 3.2% among men aged 30 to 44. In Liverpool, the estimated prevalence was 2.4% among men aged 15 to 29 and 2.8% among women aged 15 to 29. In London, the estimated prevalence was 1.5% among men aged 15 to 29 and 1.5% among women aged 15 to 29.

The London estimate, when divided into inner and outer London, was 1.2% (95% CI 1.0% to 1.5%). The estimate for inner London was 1.2% (95% CI 1.0% to 1.5%) and the estimate for outer London was 1.2% (95% CI 1.0% to 1.5%).

The estimated prevalence of injecting drug users varied by age and gender in all three cities. In Brighton, the estimated prevalence was 2.4% among men aged 15 to 29, 3.2% among women aged 15 to 29, and 3.2% among men aged 30 to 44. In Liverpool, the estimated prevalence was 2.4% among men aged 15 to 29 and 2.8% among women aged 15 to 29. In London, the estimated prevalence was 1.5% among men aged 15 to 29 and 1.5% among women aged 15 to 29.

The London estimate, when divided into inner and outer London, was 1.2% (95% CI 1.0% to 1.5%). The estimate for inner London was 1.2% (95% CI 1.0% to 1.5%) and the estimate for outer London was 1.2% (95% CI 1.0% to 1.5%).

The estimated prevalence of injecting drug users varied by age and gender in all three cities. In Brighton, the estimated prevalence was 2.4% among men aged 15 to 29, 3.2% among women aged 15 to 29, and 3.2% among men aged 30 to 44. In Liverpool, the estimated prevalence was 2.4% among men aged 15 to 29 and 2.8% among women aged 15 to 29. In London, the estimated prevalence was 1.5% among men aged 15 to 29 and 1.5% among women aged 15 to 29.

The London estimate, when divided into inner and outer London, was 1.2% (95% CI 1.0% to 1.5%). The estimate for inner London was 1.2% (95% CI 1.0% to 1.5%) and the estimate for outer London was 1.2% (95% CI 1.0% to 1.5%).

The estimated prevalence of injecting drug users varied by age and gender in all three cities. In Brighton, the estimated prevalence was 2.4% among men aged 15 to 29, 3.2% among women aged 15 to 29, and 3.2% among men aged 30 to 44. In Liverpool, the estimated prevalence was 2.4% among men aged 15 to 29 and 2.8% among women aged 15 to 29. In London, the estimated prevalence was 1.5% among men aged 15 to 29 and 1.5% among women aged 15 to 29.

The London estimate, when divided into inner and outer London, was 1.2% (95% CI 1.0% to 1.5%). The estimate for inner London was 1.2% (95% CI 1.0% to 1.5%) and the estimate for outer London was 1.2% (95% CI 1.0% to 1.5%).
men aged 15–29 in both cities. Overall in the 12 London boroughs the estimated prevalence of IDU was 1.9% among men and 0.6% among women aged 15 to 44.

Table 3 shows estimates of public health indicators associated with injecting drug use in Brighton, Liverpool, and London as a whole. The number of IDU that receive treatment annually in general practice or at a specialist drug agency was not readily available and drawn from a variety of sources. All provide convergent evidence that the proportion of IDUs receiving treatment is low: at 22% in Liverpool and 16% in Brighton.

Table 3 Public health indicators using the estimates for injectors age 15 to 44 years in Brighton, Liverpool, and London 2000/01

<table>
<thead>
<tr>
<th>Public health indicator</th>
<th>Number of events</th>
<th>Indicator</th>
<th>Number of events</th>
<th>Indicator</th>
<th>Number of events</th>
<th>Indicator</th>
</tr>
</thead>
<tbody>
<tr>
<td>Proportion of injectors receiving structured treatment†</td>
<td>363</td>
<td>16%</td>
<td>654</td>
<td>22%</td>
<td>7000</td>
<td>22%</td>
</tr>
<tr>
<td>Annual number of syringes distributed per IDU per year‡</td>
<td>429000</td>
<td>186</td>
<td>566500</td>
<td>195</td>
<td>4910000</td>
<td>143</td>
</tr>
<tr>
<td>(coverage per injection)</td>
<td>(27%)</td>
<td>(28%)</td>
<td>(20%)</td>
<td>Opioid overdose mortality rate*</td>
<td>48</td>
<td>2.1%</td>
</tr>
</tbody>
</table>

*Estimate of number of injectors based on applying prevalence estimates for inner and outer London from 12 London boroughs to all of greater London. †Brighton: number in specialist drug treatment and primary care Brighton Drug Commissioning (personal communication); London: number in specialist drug treatment based on number and proportion receiving substitute treatment reported to London's National Drug Treatment Monitoring System supplemented by an estimate of number in treatment in primary care (Dr Chris Ford, personal communication); Liverpool: local drug treatment monitoring system. ‡Brighton: Brighton Drug Commissioning (personal communication); London: 1997 Survey and local SEP data*; Liverpool: Local SEP surveillance (John Moores University). From Office of National Statistics Drug Related Deaths Database.
The estimated annual opioid mortality rate based on the number of opioid overdose deaths identified by Office of National Statistics as the numerator and the prevalence estimates as the denominator was 2.1% in Brighton compared with about 1.0% in Liverpool and 0.7% London.

DISCUSSION

These estimates indicate substantial numbers of injecting drug users in the three cities (Brighton, Liverpool, and London) and the subsequent measures of coverage serve to emphasise the need for further public health action to treat and prevent the harms associated with injecting drug use.

Credibility of the estimates

The findings suggest that between 1 in 50 and 1 in 80 of the adult population aged 15–44 in the three cities is an injecting drug user, which equates to between 10 and 18 patients in a typical general practice list of 2000 patients with 900 aged 15–44. Thus, in Brighton, Liverpool, and London the prevalence of injecting drug use among young adults is as common as diabetes and greater than many other chronic conditions such as epilepsy or psychosis. The prevalence also was higher among males and in the “older” age-group 30–44, which implies that a large proportion of the injecting population is an aging cohort.

Given abbreviated data on the subjects (initials rather than full name), there is potential for under-matching, which would lead to over-estimates of the population. Furthermore, the inadequacy of datasets in London to cover the whole of the geographical area and the information on area of residence limited analyses for inner and outer London and separate boroughs. However, these estimates are plausible. Firstly, there is a consistency in the size of the estimates, and distributions by age group and gender between the three cities. Secondly, the ratio between the unobserved and observed number of injectors were not excessive. Moreover, the number of truly hidden IDUs will be lower than the unobserved population. For instance in London, data were not collected from five local syringe exchange programmes (SEP) and three hospital Accident and Emergency departments or from any pharmacy offering syringe exchange, drug outreach team, or drop in service. Thirdly, a prevalence of over 1% has been recorded previously in Glasgow; and the higher prevalence in Brighton fits with evidence on overdose mortality. Finally, in Brighton and Liverpool where consultation has already taken place, local specialists and policy makers gave their support to the estimates.

Coverage and public health implications

The government aims to double the number of problem drug users in treatment.1 In the three sites there is ample opportunity for this, given that less than one in four IDUs are in receipt of treatment at any one time. Unfortunately data on the number in treatment were of poor quality and requires urgent improvement.

The estimated coverage of syringe distribution at approximately one clean syringe per injector every two to 2.5 days (or 20% to 27% of all injections) was similar in the three areas, adding support to the prevalence estimates. While policy makers in England deserve credit for reaching such a high coverage in comparison with many other cities and countries worldwide it should still be regarded as insufficient. Corroboration of the coverage in London especially is required. The prevalence of hepatitis C virus infection among IDUs in London, Liverpool, and Brighton is higher than many other areas in England and Wales; and the sharing of injecting equipment continues at high levels. Syringe exchange distribution needs to be expanded, perhaps doubled, to reduce the opportunity for sharing and minimise the risk of viral transmission; especially hepatitis C infection.

Annually injecting drug users are estimated generally to have a risk of fatal overdose of nearly 1%, as found for the estimates for London and Liverpool, which tends to increase with duration of injecting. In Brighton, not only was the prevalence of injecting estimated to be higher than the other cities, but the proportion of opioid users reported as dying from overdose was also estimated to be higher at 2%. It seems that Brighton may have proportionately more opioid overdose deaths than any other city in the UK. In 2000, Brighton recorded 0.4 opioid overdose deaths per 1000 adult population aged 15 to 44, compared with 0.1 in Liverpool and London, which cannot be explained entirely by differences in the size of the IDU population. Clearly, local policymakers should consider expanding the availability of substitution treatment, and continue investigating why the risk of fatal opioid overdose is so high in Brighton.

Methods and future work

Indirect methods of estimating prevalence are inherently uncertain, and the figures they produce need to be treated cautiously. There are no “simple rules” for determining reliability or the sample size required, though in general the larger the study the better the evidence. None the less, we show that by using a consistent approach and new statistical techniques (that have not been used before for drug use or multiple data sources), useful estimates describing the extent of drug use in the population can be obtained. Further work will be conducted on model selection and on exploring the limits of the number of covariates that can be included in capture-recapture analyses. Capture-recapture also has been suggested to provide the most accurate estimate of prevalence when compared with other epidemiological counting methods.

During the study, substantial if not most effort and time was spent on gaining access and collecting routine data. It is vital that prevalence estimation—through capture-recapture—should be included as a key objective of current and future data sources, such as monitoring systems for drug treatment, and the National Arrest Referral scheme (that has been suggested for the surveillance of other chronic diseases problems). Better evidence will be available therefore to support or to counter claims that insufficient numbers of injecting drug users are being treated, and to bring some rationale to debate on drug policy.

ACKNOWLEDGEMENTS

We thank the numerous individuals and services in Brighton, Liverpool, and London that allowed us to collect their data.

Authors’ affiliations

M Hickman, V Hope, Centre for Research on Drugs and Health Behaviour, Social Science and Medicine, Primary Care and Population Sciences, Imperial College, London, UK

V Higgins, M Bellis, Centre for Public Health, Liverpool John Moores University, Liverpool, UK

K Tilling, Social Medicine, University of Bristol, Bristol, UK

A Walker, Brighton and Hove Drug Action Team, Brighton, UK

J Henry, Academic Department of Accident and Emergency Medicine, Imperial College, London, UK

Funding: the study was funded by the Home Office Research and Statistics Directorate. Matthew Hickman is funded through a Department of Health Public Health Career Scientist award. The Centre for Research on Drugs and Health Behaviour is core funded by the Department of Health.

Conflicts of interest: none declared.

www.jech.com
REFERENCES


3 Davies N. How Britain is losing the drugs war. Guardian 2003;22 May.


