BMJ OPEN Prevalence of, and risk factors for, HIV, hepatitis B and C infections among men who inject image and performance enhancing drugs: a cross-sectional study

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ABSTRACT

Objective: To describe drug use, sexual risks and the prevalence of blood-borne viral infections among men who inject image and performance enhancing drugs (IPEDs).

Design: A voluntary unlinked-anonymous crosssectional biobehavioural survey.

Setting: 19 needle and syringe programmes across England and Wales.

Participants: 395 men who had injected IPEDs.

Results: Of the participants (median age 28 years), 36% had used IPEDs for <5 years. Anabolic steroids (86%), growth hormone (32%) and human chorionic gonadotropin (16%) were most frequently injected. with 88% injecting intramuscularly and 39% subcutaneously. Two-thirds also used IPEDs orally. Recent psychoactive drug use was common (46% cocaine, 12% amphetamine), 5% had ever injected a psychoactive drug and 9% had shared injecting equipment. 'Viagra/Cialis' was used by 7%, with 89% reporting anal/vaginal sex in the preceding year (20% had 5+ female-partners, 3% male-partners) and 13% always using condoms. Overall, 1.5% had HIV, 9% had antibodies to the hepatitis B core antigen (anti-HBc) and 5% to hepatitis C (anti-HCV). In multivariate analysis, having HIV was associated with: seeking advice from a sexual health clinic; having had an injection site abscess/wound; and having male partners. After excluding those reporting male partners or injecting psychoactive drugs, 0.8% had HIV, 8% anti-HBc and 5% anti-HCV. Only 23% reported uptake of the hepatitis B vaccine, and diagnostic testing uptake was poor (31% for HIV, 22% for hepatitis C).

Conclusions: Previous prevalence studies had not found HIV among IPED injectors. HIV prevalence in this, the largest study of blood-borne viruses among IPED injectors, was similar to that among injectors of psychoactive drugs. Findings indicate a need for targeted interventions.

ARTICLE SUMMARY

Article focus

- The Vulnerability to infection of people who inject drugs is widely recognised; however, studies have rarely focused on users of image and performance enhancing drugs. These drugs can be used to change one's appearance for aesthetic reasons, as well as to improve performance.
- Over the last decade, the number of men using needle and syringe programmes who report injecting image and performance enhancing drugs has risen in England and Wales; as a result, there has been increased concern about the levels of bloodborne viral infections in this group.
- This study describes the nature of drug use and the risk behaviours in this population, as well as the prevalence of HIV, hepatitis B and C.

Key messages

- The overall prevalence of HIV among men injecting image and performance enhancing drugs was similar to that among those injecting psychoactive drugs in England and Wales. Previous prevalence studies of people who inject image and performance enhancing drugs had not detected HIV.
- When the results of this study are compared with those of a previous study undertaken in England and Wales in the mid-1990s, they suggest that the prevalence of hepatitis B infection among injectors of image and performance enhancing drugs *might* have increased over time.
- Sexual risk behaviours and psychoactive drug use were common among injectors of image and performance enhancing drugs, and the sharing of injecting equipment was also reported. The uptake of diagnostic testing for blood borne-viral infections and the vaccine against hepatitis B was low.

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ARTICLE SUMMARY

Strengths and limitations of this study

- This study recruited image and performance enhancing drug users through needle and syringe programmes. Injectors of these drugs who are not in contact with these services may have a different risk profile and levels of infection.
- Oral-fluid testing was used to detect antibodies to HIV, hepatitis B and C; however, tests on these samples for anti-HCV and anti-HBc have reduced sensitivity.
- This is the largest study of blood-borne viruses among men who inject image and performance enhancing drugs; however, the sample size still restricts its power. Consequently, caution is needed when attempting to generalise these findings.

INTRODUCTION

The vulnerability of people who inject drugs (PWID) to HIV and other infections is widely recognised; however, studies have focused on individuals who inject psychoactive drugs (such as opiates and stimulants) rather than on those who inject drugs to enhance image and performance.^{1–4} The number of injectors of image and performance enhancing drugs (IPEDs) in contact with needle and syringe programmes (NSPs) has grown substantially in the UK,⁵ and there has been increasing concern about the use of IPEDs and the associated harms in the UK and elsewhere.^{3 5–11}

A range of illicit drugs can be injected with the aim of changing image and performance. These drugs range from tanning drugs, such as 'Melanotan-II',¹² to those used in body-building, such as human growth hormone.^{3 13} The most commonly injected and studied IPEDs are anabolic steroids (AS).^{3 5} IPEDs are taken both orally and by injection, with some being predominantly injected and others being taken only orally. Many users of these substances also take an array of different drugs.^{3 5 14} The use, and particularly the injection of IPEDs has been associated with a range of harms including infections caused by bacteria^{15–19} and blood-borne viruses (BBVs).^{6 20–25}

In England and Wales (E&W), surveillance of HIV and viral hepatitis among PWID is undertaken through an annual unlinked-anonymous survey^{26 27} targeted at injectors of psychoactive drugs. A very small number of IPED injectors participated in this survey²⁰; among the 149 sampled during the 1990s, 2% had antibodies to the hepatitis B core antigen (anti-HBc, a marker of having ever been infected with hepatitis B virus (HBV))) and none had antibodies to HIV (anti-HIV).²⁰ In a surveillance study of NSP clients in Australia, 1.6% (n=318) of those participating over a 10-year period reported steroid injection, with 10% having antibodies to hepatitis C virus (anti-HCV) and none having anti-HIV.²¹ Only one other survey of IPED injectors has collected biological samples; this study purposively recruited 63 AS injectors in Victoria, Australia and found that 12% had anti-HBc, 9.5% anti-HCV and none anti-HIV.⁶ A second

Australian study found that half of the IPED users sampled had ever experienced an injection-related health problem, with 6% having ever had an abscess.⁸

A number of other UK studies have recruited IPED injectors, principally AS injectors; however, none of these collected biological samples. These studies were mostly small (N<100), and typically recruited through gyms,^{28–33} with two recruiting gay men.^{34 35} The prevalence of ever sharing injecting equipment in these studies ranged from 0.3% to 6%,^{20 28–30 32} but in one study it was 20%.³³ The sharing of drug vials was more common (2.4%³⁵; 9.9%³⁴; 23%³²). Studies elsewhere have found similar levels of equipment sharing.^{6 8} IPED users also report using psychoactive drugs, particularly stimulants, though the reported injection of psychoactive drugs is rare.^{6 8 28 35 36} IPED users also tend to have more sexual partners than their comparison groups^{20 28} and report risky sexual behaviours^{20 32} and low levels of condom use,^{28 34} suggesting an elevated risk for HIV infection through sexual activity.

During 2010 and 2011, in response to the increasing concerns about IPED use, a targeted survey was undertaken as part of the ongoing unlinked-anonymous survey of PWID. The aim of this survey was to describe the: (1) patterns of drug use and injecting risk; (2) sexual behaviours and (3) BBV prevalence among IPED injectors. As far as we are aware, this is the largest study, and the first outside Australia, to purposively recruit IPED injectors to measure the prevalence of anti-HIV, anti-HBc and anti-HCV.

METHODS

Recruitment

In E&W, PWID have been recruited into a voluntary unlinked-anonymous monitoring survey since 1990, and the methodological details of this cross-sectional survey have been published previously.^{26 27 37} Briefly, agencies providing services to PWID (eg, NSPs and addiction treatment) at sentinel locations throughout E&W invite clients who have ever injected to participate. Sentinel sites are selected so as to reflect the geographic distribution and range of services offered to PWID. Those who consent to participate (overall refusal rate during 2010/ 2011, 4.7%) provide a biological sample and selfcomplete a brief questionnaire focused on psychoactive drug use.^{26 27 37} The survey has multisite ethics approval. This study purposively recruited IPED injectors through 19 sites that provided NSPs. Participants were recruited either when attending an NSP site or through outreach provision; they provided an oral-fluid sample and selfcompleted a short, specially developed, questionnaire focused on IPED use (types of drug used and routes of administration), related behaviours (injecting practices and sexual behaviours) and health service use.

Laboratory methods

Oral-fluid specimens were collected using the OraSure device (OraSure Technologies Inc, Pennsylvania, USA).

These were tested for anti-HIV using an in-house GACELISA with similar performance to GACELISA HIV 1+2 (Abbott Murex Diagnostics Ltd, Dartford, UK). Reactive specimens underwent further testing according to a proven algorithm that included a second ELISA and Western blot (sensitivity and specificity approaches 100%³⁸). Anti-HCV testing employed a previously validated commercial enzyme-immunoassay (Ortho HCV 3.0 SAVe, Ortho Diagnostics) with 92% sensitivity and 99% specificity,³⁹ and for anti-HBc an in-house IgG classspecific antibody capture EIA procedure was used which had an estimated sensitivity of 75% and specificity of 99% (JV Parry and A Judd, personal communication). The oral-fluid sample quality was verified by testing each sample for the presence of a predetermined minimum quantity of total IgG (1 mg/L) employing an in-house ELISA method.

Analyses

Descriptive analyses were first undertaken, and then bivariate associations (p<0.05) between outcome variables (anti-HIV, anti-HBc and anti-HCV positivity, equipment sharing and condom use) and covariates (age, drug use, sexual practice and health services use; table 1) were examined using Fisher's exact (when expected cell frequencies <5) and Pearson's χ^2 tests. Where possible associations were found (p<0.10), these were further examined through logistic regression models using the forward stepwise procedures to select variables, with selection based on the likelihood ratio test (p<0.05). All analyses were undertaken using SPSS V.19.

RESULTS

Between May 2010 and May 2011, 400 IPED injectors participated in this study; five (1.25%) women were excluded from the analyses (due to the small number). The participants' characteristics and health service usage are summarised in table 1. Of those reporting their age (88%, n=347), a quarter (27%) were aged <25 years. During the preceding year, 45% had seen a general practitioner and 28% had taken the prescribed medication.

Drug use

Details of the participants' IPED use during the preceding year are given in table 1. AS were the most commonly injected IPED (86%). Over half of the participants reported consuming these orally (57%), a third reported injecting growth hormone (32%) and almost a quarter reported using oral antioestrogens (23%). Overall, 65% (n=252) had taken an IPED orally during the preceding year, with 58 (23%) of these having taken two types orally, and 85 (34%) \geq 3 types. Most had injected only one type of IPED during the preceding year; however, 87 (22%) had injected two types and 58 (15%) \geq 3 types. Considering injecting and oral use, 71 (18%) had taken two types of IPED and 133 (34%) \geq 3 during the preceding year. Those who injected human growth hormone were more likely to be older (aged >35 years) than those who had not $(37\% \ (47/128) \ vs \ 22\% \ (60/267), \ p<0.001)$; there were no other significant differences in the IPEDs used by age. During the preceding year, most of the participants (74%) reported that they had usually injected themselves, and the majority (88%) had injected intramuscularly (table 1).

The participants also reported psychoactive drug use (table 1), with 46% snorting cocaine and 12% snorting, drinking or swallowing amphetamine during the preceding year. Ever having injected a psychoactive drug (including heroin and cocaine) was reported by 4.8% (table 1). Those who had injected a psychoactive drug were more likely to report injecting insulin as an IPED than those who had not (21% (4/19) vs 4.8% (18/376), p=0.016); there were no other significant differences in the IPEDs used between those who had injected psychoactive drugs and those who had not.

Overall, 8.9% (95% CI 6.4% to 12%) reported having ever shared a needle/syringe or drugs vial (table 1); 27 (6.8%) had just shared a vial, 6 (1.5%) had just shared a needle/syringe and 2 (0.51%) had shared both. Factors associated with sharing are summarised in table 2. In the multivariable analysis, ever having shared a needle/ syringe or drug vial was associated with having ever injected a psychoactive drug, having sought advice from an sexual health/sexual transmitted infections (SH/ STI) clinic, subcutaneous injection and having snorted, drunk or swallowed amphetamine (table 2).

Sexual behaviour

Nine-tenths (89%, 350/395) reported having anal or vaginal sex in the preceding year, and 9.1% (36/395) had ≥ 10 partners (table 1). Considering just female partners, 20% (80/395) of respondents had \geq 5 partners. Thirteen (3.3%) reported ≥ 1 male sexual partner during the preceding year (table 1). Those reporting male sexual partners were older than those who did not (median age 38 years, IQR 12; and 28 years, IQR 11, respectively). Those reporting male sexual partners were also more likely to have ever injected a psychoactive drug (23% (3/13) vs 4.2% (16/382), p=0.020), more likely to report snorting, drinking or swallowing amphetamine during the last year (46% (6/13) vs 11%(41/382), p=0.002), and a higher proportion reported snorting cocaine, but this was not significant (62%) (8/ 13) vs 45% (173/382), p=0.248). Those reporting male sexual partners were also more likely to report having ever shared a needle/syringe or vial (25% (4/13) vs)8.1% (31/382), p=0.021). A higher proportion of those reporting male sexual partners reported always using condoms during the last year, but this difference was not significant (38% (5/13) vs 19% (73/382), p=0.146). There were no differences in the types of IPED used, nor in their routes of administration, between those reporting male partners and those not.

Characteristic			n
Demographic			
Age, years	Under25	27%	106
	25-34	34%	134
	35 and over	27%	107
	Median (IQR)	28 (13	
	Not reported	12%	 48
Had ever been in prison		16%	63
Health service use			
Had ever used a Needle and Syringe Programme		75%	298
Had seen a general practitioner in the last year about their health		45%	178
Had you got advice at an Accident & Emergency / Walk-in in the last ye	ar	16%	64
Had taken/used prescribed medication in last year		28%	111
Had sought advice from a sexual health / sexually transmitted infections	s clinic in the last year	17%	68
Had been vaccinated against hepatitis B	,	23%	90
Had had a blood test for hepatitis C		22%	85
Had had a blood test for HIV		31%	122
Symptom of injury or infection at injection site		01/0	
Had ever had redness at an injection site		43%	168
Had ever had an injection site abscess/sore/open wound		6.80%	27
Image and performance enhancing drug use, last year		0.0070	_,
Years since first used an image and performance enhancing drug	0-4	36%	141
reals since met deed an image and performance ermanoling drug	5+	32%	128
	Median (IQR)	4 (8)	120
	Not reported	32%	126
Oral anabolic steroids	Not reported	57%	226
Oral anti-oestrogens		23%	92
Oral clenbuterol		15%	92 60
Oral ephedrine		20%	78
Oral thyroid hormones		20 % 9%	37
Oral phosphodiesterase type 5 inhibitor (PDE5i; "Viagra/Calias")		9 % 6.6%	26
Oral other image and performance enhancing drug (inc. Diuretics,		12%	46
		12/0	40
2,4–dinitrophenol and Pro/designer) Injected anabolic steroids		86%	340
Injected growth hormone		32%	128
Injected human chorionic gonadotropin (hCG)		32 % 16%	62
		5.6%	22
Injected insulin injected Injected melanotan		5.6% 8.6%	22 34
•	'n	5.1%	20
Injected other image and performance enhancing drug (inc. ethryopoeti	11,	5.1%	20
insulin–like growth factor 1 and nalbuphine hydrochloride) Use of other illicit drugs			
Ever injected illicit drug other than an image and performance enhancir	a drug	4.8%	19
Snorted cocaine in the last year	ig ulug.	4.8%	181
Shorted, drunk or swallowed amphetamine in last year		40 % 12%	47
		12%	47
Injecting practice		170/	60
Who usually injected you, last year?	Someone else	17% 74%	68
	Myself		294
Intromuce user intertion in the last year	Not reported	8.4%	33
Intramuscular injection in the last year		88%	346
Subcutaneous injection in the last year		39%	154
Ever shared needle, syringe or vial		8.9%	35
Sexual behaviour	0.14	000/	450
Number of sexual partners last year	One	38%	152
	Two or more	47%	187
	Not reported/no sex	14%	56
Gender of sexual partners last year	Male partner(s)	3.3%	13
	No male partners	82%	323
	Not reported/no sex	15%	59
Always condom (anal/vaginal sex) or no sex last year		20%	78

 Table 2
 Factors associated with risk behaviours among the IPEDs sampled

Table 2 Factors associated with risk behavio	Total	Yes		Unadjusted odds ratio with 95% Cl				Adju with		atio	
Ever shared a needle, syringe or vial	395	35	8.9%								
Number of sexual partners in the last year											
One	152	8	5.3%	0.3	0.12	-	0.94	*			
Two or more	187	19	10%	0.7	0.28	-	1.6				
Not reported/no sex	56	8	14%	1.0							
Pearson χ^2 test	p=	0.088									
Gender of sexual partners in the last year											
Male partner(s)	13	4	31%	5.5	1.6	-	19	*			
No male partners	323	24	7.4%	1.0							
Not reported/no sex	59	7	12%	1.7	0.69	-	4.1				
Pearson χ^2 test	p=	0.010									
Injected illicit drugs other than IPED											
Yes	19	7	37%	7.2	2.6	-	20	6.3	2.1	-	19
No/not reported	376	28	7.4%	1.0				1.0			
Fisher's exact test	p=	0.001									
Have you got advice from a SH/STI clinic in th	-						_				
Yes	68	11	16%	2.4	1.1	-	5.2	2.2	1.0	-	5.1
No/not sure	327	24	7.3%	1.0				1.0			
Pearson χ^2 test	p=	0.020									
Injected growth hormone (as IPED)											
Yes	128	18	14%	2.4	1.2	-	4.8	*			
No	267	17	6.4%	1.0							
Pearson χ^2 test	p=	0.012									
Injected Insulin (as IPED)		-	000/					+			
Yes	22	5	23%	3.4	1.2	-	9.8	â			
No Fisharia avaat taat	373	30	8.0%	1.0							
Fisher's exact test	p=	0.035									
Subcutaneous injection in the last year? Yes	154	21	14%	2.6	1.3		5.2	3.0	1.4		6.5
	154 241	21 14			1.3	-	5.2		1.4	_	0.5
No/not sure Pearson χ^2 test		0.008	6%	1.0				1.0			
Snorted cocaine in the last year?	p=	0.000									
Yes	181	22	12%	2.1	1.0	_	4.4	*			
No	214	13	6.1%	1.0	1.0	_	4.4				
Pearson χ^2 test	D=	0.034	0.170	1.0							
Snorted, drunk or swallowed amphetamine in t											
Yes	47	11	23%	4.1	1.9	_	9.1	4.1	1.7	_	9.8
No	348	24	6.9%	1.0	1.5		0.1	1.0	1.7		0.0
Pearson χ^2 test	р=	0.0002	0.070	1.0				1.0			
Always used condom for anal/vaginal sex Gender of sexual partners in the last year	350	48	14%								
Male partner(s)	13	5	38%	8.1	0.8	-	83	14	1.3	-	155
No male partners	323	42	13%	1.9	0.25	-	15	2.8	0.35	-	22
Not reported	14	1	7.1%	1.0				1.0			
Pearson χ^2 test	<i>p</i> =	0.025									
Have you ever had a blood test for hepatitis C											
Yes	82	17	21%	3.2	1.1	-	9	*			
No	201	26	13%	1.8	0.68	-	5.0				
Not sure	67	5	7.5%	1.0							
Pearson χ^2 test	p=	0.057									
Injected anabolic steroids											1.5
Yes	304	46	15%	3.9	0.92	-	17	4.2	0.96	-	18
No	46	2	4.3%	1.0				1.0			
Pearson χ^2 test	<i>p=</i>	0.048									
										Coi	ntinued

Table 2 Continued

	Total	•				-			isted oc 95% Cl		atio
Intramuscular injection in the last year?											
Yes	310	46	15%	3.3	0.77	-	14	*			
No/not sure	40	2	5.0%	1.0							
Pearson χ^2 test	p=	0.089									
Snorted cocaine in the last year?											
Yes	162	11	7%	0.3	0.1	-	0.6	0.2	0.12	-	0.52
No	188	37	20%	1.0				1.0			
Pearson χ^2 test	p=	0.0005									
*Not in final model.											

IPED, image and performance enhancing drugs; SH/STI, sexual health or sexual transmitted infections clinic.

Among those who reported sex during the preceding year, 14% (95% CI 11% to 18%, 48/350) had always used condoms. Factors associated with condom use are given in table 2. Always using condoms among those who had had sex during the preceding year was associated in the multivariable analysis with having had a male sexual partner and having not snorted cocaine (table 2).

BBV prevalence

Overall, 1.5% (95% CI 0.7% to 3.3%; n=6) had anti-HIV, 8.8% (95% CI 6.4% to 12%) had ever been infected with hepatitis B (26 anti-HBc positive, adjusted for test sensitivity of 0.75) and 5.5% (95% CI 3.7% to 8.2%) with hepatitis C (20 anti-HCV positive, adjusted for test sensitivity of 0.92). Covariates associated with anti-HIV, anti-HBc or anti-HCV positivity are given in table 3. In the multivariable analysis, anti-HIV positivity was associated with having male sexual partners in the preceding year, ever having an abscess/sore/open wound at the injection site, and having sought advice from an SH/STI clinic in the preceding year (table 3). Having anti-HBc was associated in the multivariable analysis with having obtained advice from an SH/STI clinic and having not injected oneself subcutaneously in the preceding year (table 3). Anti-HCV positivity was associated with having ever injected a psychoactive drug and having taken a phosphodiesterase type 5 inhibitor (PDE5i) in the preceding year in the multivariable analysis (table 3).

After excluding those who reported either sex with men or having ever injected a psychoactive drug, 0.8% had anti-HIV (95% CI 0.28% to 2.4%, 3/366), 8.0% anti-HBc (95% CI 5.6% to 11%, adjusted for test sensitivity, 22/366) and 4.7% anti-HCV (95% CI 2.9% to 7.3%, adjusted for test sensitivity, 16/366), with 10% (95% CI 7.7% to 14%, 38/366) having one or more of these three markers. In this group, having anti-HIV was found to be associated only with ever having had an abscess/wound at an injection site (8% (2/25) vs 0.29% (1/341) for those who had not, p=0.013), and having anti-HBc was only associated with having sought advice from an SH/STI clinic in the preceding year (16% (9/56) vs 4.2% (13/310) for those who had not, p=0.002). The use of three types of IPEDs was associated with having anti-HCV: having taken a PDE5i (21% (5/24) vs 3.2% (11/342) for those who had not, p=0.002); having injected insulin as an IPED (18% (3/17) vs 3.7% (13/349) for those who had not, p=0.032); and having injected a less commonly used IPED (17% (3/18) vs 3.7% (13/348) for those who had not, p=0.037).

In total, 47 (12%, 95% CI 9.1% to 15%) were positive for one or more of the anti-HIV, anti-HBc and anti-HCV, with 43 having just one of these markers and four having two or more of these markers. Two had anti-HBc and anti-HCV, one of these reported injected psychoactive drugs, and neither reported having had sex with men. One, who reported sex with men and injecting psychoactive drugs, had anti-HIV and anti-HBc; the remaining participant had all three markers and did not report either sex with men or injecting psychoactive drugs.

Uptake of interventions related to the three BBV infections was poor. Overall, only 23% reported receiving a dose of HBV vaccine, 31% ever having a diagnostic HIV test and 22% having an HCV test (table 1).

DISCUSSION

IPED injectors are at risk of infection with HIV as well as other BBVs. This is the first prevalence study to have found HIV among IPED injectors, with the prevalence at 1.5%, similar to that found among injectors of psychoactive drugs in England and Wales (1.2%, 2011^{40}). However, anti-HBc and anti-HCV—at 8.8% and 5.5%, respectively—are lower than among psychoactive drug injectors (16% and 43%, respectively, 2011^{40}). The prevalence of all three BBVs in this sample would appear to be higher than that found in the general UK population.⁴¹ ⁴² Once those who reported either sex with men or injecting psychoactive drugs were excluded, 10% had been infected with one or more of HIV, hepatitis B and C.

It is important to consider the limitations of this study. The comparative rarity, marginalisation and illicit nature of injecting drug use impede the construction of a sampling frame, making the representativeness of our sample impossible to measure. This study used an established methodology for recruiting PWID through specialist services²⁶ ²⁷; however, the robustness of this

Table 3 Factors associated the blood-borne virus infections among the IPED	s sampled
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Total Positive ratio with 95% CI with 95% CI HV 396 6 1.5% Age, yars' Age, yars' 0.3% 1.0 -<	Table 3 Factors associated the blood-borne virus infections among the IPEDs sampled Unadjusted odds Adjusted odds ratio													
Age: Age: Single unknown 287 1 0.3% 1.0 '		Total	tal Positive							-				
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Aged 35 or age unknown 27 1 0.3% 1.0 - 122 - - - 121 - - - - 121 - - - 13 3 23% 38 6.79 - 211 - - 13 3 23% 38 6.79 - 211 - - 145 - 15 - 15 - 15 - 10 - <		000	0	1.570										
Aged 35 or over 102 5 4.9% 14 1.62 - 122 Fisher's exact test p= 0.006 3 3 23% 38 6.79 - 211 *<		287	1	0.3%	1.0				*					
Fisher's exact test p= 0.006 Gender of sexual partner in the last year's 382 3 0.8% 1 - - 211 * - - 1 - - - 11 * - - 11 * - - 10 - - 13 3 0.8% 1 0 - - 10 - - 10 - 11 - 10 - 11 - 10 - 11 - 10 - 11 - 10 - 11 - 10 - 179 - 100 - 179 - 100 - 10 - 10 -						1.62	_	122						
Gender of sexual partners in the last year* 1 3 3 2.3% 38 6.79 - 211 *	•				••									
Maile sexual partmer No male partmers in the last year and age (in years)' Male sexual partmers, aged x35 or age not reported Discretal distribution of the partmers, aged x35 or age not reported Paramonic for the partmers, partmers, aged x35 or age not reported Paramonic for the partmers, part		P												
No male pathmerino sex 32 3 0.8% 1 Fisher's exact test p= 0.001 Gender of sexual pathmers in the last year and age (in years)* 3 3 2.2% 85 8.13 - 893 79 4.29 - 145 No Male pathners, aged 35 or age not reported 265 1 0.4% 1.0		13	3	23%	38	6.79	_	211	*					
Fisher's exact test $p=$ 0.001Gender of sourcal partners in the last year and age (in years)*13323%858.13 $-$ 893794.29 $-$ 145No Male partners, aged <35 or age not reported				0.8%	1									
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		241	20	8.3%	2.2	0.88	-	5.7	2.6	0.99	-	6.7		
Continue	Pearson χ^2 test	p=	0.085											
Continue											Со	ntinued		

Table 3 Continued

	Total	Positiv	/e		justed with 95				isted 95%		ratio
Anti-HCV	395	20	5.1%								
Age, years											
Under 25	106	4	3.8%	1.0				†			
25–34	134	3	2.2%	0.58	0.13	—	2.7				
35 and over	107	10	9.3%	2.6	0.80	-	8.7				
Not Reported	48	3	6.3%	1.7	0.37	-	7.9				
Pearson χ^2 test	p=	0.078									
Injected illicit drugs other than IPED											
Yes	19	3	15.8%	4.0	1.1	-	15	4.4	1.1	-	17.2
No/not Sure	376	17	4.5%	1.0				1.0			
Fisher's exact test	p=	0.064									
Taken or used any prescribed medication in the last	t year?										
Yes	111	10	9.0%	2.7	1.1	-	7	†			
No/not sure	284	10	3.5%	1.0							
Pearson χ^2 test	p=	0.025									
Have you ever had a blood test for hepatitis C?											
Yes	85	8	9.4%	1.7	0.55	-	5.6	†			
No	221	7	3.2%	0.5	0.17	-	1.8				
Not sure	89	5	5.6%	1.0							
Pearson χ^2 test	p=	0.080									
Taken PDE5i: Viagra/cialis											
Yes	26	5	19%	5.6	1.9	-	17	6.0	1.9	-	18
No	369	15	4.1%	1.0				1.0			
Fisher's exact test	p=	0.007									
Injected insulin (as IPED)											
Yes	22	4	18%	5.0	1.5	-	16	†			
No	373	16	4.3%	1.0							
Fisher's exact test	p=	0.019									
Other injected IPED (inc. EPO, IGF-1 and Nubain)											
Yes	20	3	15%	3.7	0.99	-	14	†			
No	375	17	4.5%	1.0							
Fisher's exact test	p=	0.073									
Ever shared a needle, syringe or vial?											
Yes	35	4	11%	2.8	0.87	-	8.8	†			
No/not sure	360	16	4%	1.0							
Fisher's exact test	p=	0.090									

*Combined variable entered into the final model due to an interaction between age and reporting male sexual partners: with 77% of those with male sexual partners aged over 35 years compared with 25% of those not reporting male sexual partners. †Not in the final model.

SH/STI, sexual health or sexual transmitted infections clinic; Anti-HCV, antibodies to the hepatitis C virus; Anti-HBc, antibodies to the hepatitis B core antigen; EPO, erythropoietin; IGF-1, insulin–like growth factor 1; IPED, image and performance enhancing drugs; PDE5i, phosphodiesterase type 5 inhibitor.

approach for IPED injectors is unknown and cannot currently be assessed due to the very limited knowledge on the size and nature of this group.^{3 5} The use of NSPs to access this group was a pragmatic approach; communitybased recruitment approaches, such as Respondent Driven Sampling and Time-Location Sampling, that are often advocated for hard to reach populations⁴³ are possible alternatives. However, these are likely to be difficult to implement with this group due to the diversity of the drugs used, the clandestine and close-knit nature of this group and because drug use usually takes place in private settings (such as homes or gyms).^{3 5} The findings here also rely on self-reported behaviours—though their reliability has not been assessed among IPED users, these have been found to be reliable for psychoactive drug injectors⁴⁴ ⁴⁵—and infection with BBVs has been determined by laboratory-based biological data from the testing of oral-fluid samples. While oral-fluid testing is highly sensitive for anti-HIV, the sensitivity is reduced for anti-HCV and anti-HBc.³⁹ While this is the largest study of BBVs in this population, the sample size still restricts its power, and consequently caution is needed when attempting to generalise these findings to the wider population of IPED injectors.

The levels of HIV and anti-HBc are higher than in the only previous UK study to measure these in IPED

injectors. Undertaken >10 years ago using a similar methodology, this found no HIV and an anti-HBc prevalence of 2%;²⁰ suggesting that the prevalence of these infections among IPED injectors might have increased over time. Exposure to BBVs among IPED injectors appears to be associated with sexual risks and the injection of psychoactive drugs; although injecting psychoactive drugs is rare among IPED injectors, unprotected sex with multiple partners is common. The sharing of injecting equipment or drug vials among IPED injectors at 8.9% is much less common than among injectors of psychoactive drugs (in 2011, 37% of psychoactive drug injectors reported recently sharing injecting equipment⁴⁰). Though exposure was not associated with sharing in the multivariate analyses in this study, BBV transmission through IPED injection cannot be excluded as this study may have lacked sufficient power to detect this.

The associations between having HIV and the use of SH/STI clinics and having male sexual partners suggest that HIV transmission among IPED injectors might be related to sexual activity. This association may reflect AS use by some HIV positive gay and bisexual men to mask the longer-term effects of HIV infection.³⁵ The association with having an abscess/wound at the injection site probably reflects the greater vulnerability of PWID with HIV to injection-related bacterial infections.46 47 The association between exposure to HBV and having obtained advice from an SH/STI clinic is again suggestive of a role for sexual risk in infection, particularly as sexual transmission is the main route by which HBV is now acquired within the UK.48 The association between having anti-HBc and not injecting subcutaneously suggests that certain patterns of IPED use might be related to increased risk, as some IPEDs are only injected subcutaneously and others only intramuscularly, and many users take several types.³ ⁸ ^{12–14} Exposure to HCV, however, would appear to be associated with the injection of psychoactive drugs, an association that has been previously noted⁶; this finding might reflect more frequent injecting in this subgroup. Sexual activity may also play a role, assuming that the use of PDE5i is related to improving or maintaining sexual performance. These associations all require further investigation.

The level sharing found here was in line with that in preinjecting studies of risk among IPED vious users.²⁰ ^{28–30} ³² ³³ ³⁵ The association between sharing and subcutaneous injection suggests that sharing-like HBV exposure-may be associated with certain patterns of IPED use. As in previous studies of IPED injectors, sexual activity was common and condom use was poor.20 28 34 Condom use was higher among those with male sexual partners and lower among those who reported snorting cocaine. The more frequent use of condoms by gay and bisexual men probably reflects an awareness of their increased HIV risk.⁴² The association with cocaine use might possibly be related to its use as a sexual stimulant, with this possibly related to attempts to counteract the reduced libido experienced on discontinuation of AS use or in the periods between

courses of AS use ('off-cycles').¹⁴ In part, this effect may be as a result of the decrease in endogenous testosterone production,⁴⁹ which is why IPED users self-treat with human chorionic gonadotrophin in an attempt to stimulate endogenous production, with PDE5i used to symptomatically treat erectile dysfunction.⁵ ¹⁴ ⁵⁰ Increased libido following AS administration is also reported by users,¹⁴ ³³ ³⁴ ⁵¹ with similar effects being reported following the use of drugs such as melanotan-II.¹²

Associations were found between psychoactive drug use and sharing and poor condom use. IPED users who also use psychoactive drugs may be a higher risk—or perhaps less risk averse—subgroup. While this needs further investigation, it suggests—considering the substantial levels of psychoactive drug use found here and in previous studies^{6 8 28 35 36}—that those using IPED and psychoactive drugs should be an important target group for harm reduction interventions.

This study indicates that those providing services to PWID—particularly NSPs, outreach services and general practitioners—should be alert to the needs of those who use IPEDs. In particular, they need to be aware of the range of drugs that may be used by this group and of the associated injecting practices, as these differ from those of psychoactive drug injectors. Considering the BBV prevalence and levels of risk found, specialist services for PWID need to engage with IPED users and ensure that they have access to appropriate injecting equipment and targeted harm reduction advice. They should also ensure that this group has access to testing for BBVs, hepatitis B vaccinations, sexual health services and condoms.

Our findings suggest that sexual risk and the use, and particularly the injection, of psychoactive drugs are possibly the most important factors associated with BBV transmission among IPED injectors. The transmission of HIV and other BBVs through the injecting of IPED cannot be excluded, and this is certainly possible as equipment sharing does occur. However, the participants in this study were largely recruited through NSPs providing injecting equipment and advice. IPED injectors not in contact with NSPs may have a different risk profile and infection risk. Even so, our findings suggest the need for targeted interventions to address sexual health needs, psychoactive drug use and the injection practices among IPED injectors. Considering the limitations of this study, a larger study recruiting from a wider range of settings and collecting dried-blood samples is needed to more fully examine prevalence and, in particular, the associated risk factors and thus the role of IPED injection in transmission of HIV and other BBVs.

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