

Sex-Related Disparities in Criminal Justice and HIV Treatment Outcomes: A Retrospective Cohort Study of HIV-Infected Inmates

Jaimie P. Meyer, MD, MS, Javier Cepeda, MPH, Faye S. Taxman, PhD, and Frederick L. Altice, MD, MA

Compared with any other country worldwide, the United States incarcerates the highest proportion of its citizens,^{1,2} especially those with underlying medical and psychiatric disorders.³ The central purpose of correctional systems is to administer punishment for crimes and protect the public by rehabilitating offenders. Treatment of inmates' perceived nonurgent medical issues is thus often deprioritized, frequently at the expense of individual and public health.^{4,5} Incarceration is most disruptive to treatment continuity for people living with HIV/AIDS (PLWHA),⁶ who are often isolated from systems of care by poverty, racial or ethnic group, and substance use. One-sixth of the nearly 1 200 000 PLWHA in the United States cycle through criminal justice (CJ) settings annually,⁷ further constraining state budgets.⁸ It is thus critical to identify PLWHA who could benefit from alternatives to incarceration that reduce medical and custodial costs.

Women with HIV are vulnerable to incarceration because of associated drug use behaviors, commercial sex work, and having high-risk male partners.^{9,10} CJ-involved women are twice as likely as CJ-involved men and 15 times as likely as community-based women to have HIV and meet AIDS-defining criteria,^{6,11} reinforcing the need to keep HIV-infected women in the community and engaged in care. Women initiating antiretroviral therapy (ART) in community settings experience higher longitudinal survival rates than men.¹² Incarceration is destabilizing, with high associated fiscal, ethical, and social costs, and it should remain the last resort to identify and treat HIV, though women are twice as likely as men to achieve viral suppression when treated in prison.^{13,14}

After prison release, ART benefits wane for both women and men.^{13,15,16} During this chaotic transition period, suboptimal engagement in care, paired with reduced ART adherence and persistence, increases risk of

Objectives. We evaluated sex-related differences in HIV and criminal justice (CJ) outcomes.

Methods. We quantified sex-related differences in criminal offenses, incarcerations, and HIV outcomes among all HIV-infected inmates on antiretroviral therapy (ART) in Connecticut (2005–2012). Computed criminogenic risk scores estimated future CJ involvement. Stacked logistic regression models with random effects identified significant correlates of HIV viral suppression on CJ entry, reflecting preceding community-based treatment.

Results. Compared with 866 HIV-infected men on ART (1619 incarcerations), 223 women (461 incarcerations) were more likely to be younger, White, and medically insured, with shorter incarceration periods (mean=196.8 vs 368.1 days), mostly for public disorder offenses. One third of both women and men had viral suppression on CJ entry, correlating positively with older age and having treated comorbidities. Entry viral suppression inversely correlated with incarceration duration for women and with criminogenic risk score for men.

Conclusions. In the largest contemporary cohort of HIV-infected inmates on ART, women's higher prevalence of nonviolent offenses and treatable comorbidities supports alternatives to incarceration strategies. Sex-specific interventions for CJ populations with HIV effectively align public health and safety goals. (*Am J Public Health.* 2015;105:1901–1910. doi:10.2105/AJPH.2015.302687)

genotypic resistance and continued HIV transmission, with negative consequences for individual and public health.^{13,15,17,18} Just 26% of men and 25% of women have viral suppression on jail entry, which reflects a lack of treatment engagement in the community.¹⁹ Women are significantly less likely than men to (1) report taking any prescribed ART immediately before jail entry,²⁰ (2) fill an ART prescription in the 60 days after prison release,²¹ or (3) be retained in longitudinal HIV care with sustained viral suppression after release from jail.^{19,22} For CJ-involved women, HIV treatment persistence may be preferentially disrupted by untreated substance use and psychiatric disorders and by partner violence.^{23–26} We have previously reported that compared with men, women are significantly more likely to experience homelessness, ongoing cocaine use, and depression immediately before jail entry and 6 months after jail release—factors

that contribute to loss of viral suppression after release.¹⁹

Aside from HIV, considerable sex disparities prevail in terms of frequency and type of criminal offending²⁷ and sentencing standards. Women receive shorter federal sentences than men when matched on charged offense and criminal history,^{28,29} particularly with regard to property and drug-related offenses.³⁰ In some states, women have longer sentences than men.³¹ Some have argued that women's pathways to crime make incarceration less persuasive,³² requiring alternatives that more holistically address substance use and psychiatric treatment needs. Although previous studies have suggested a direct correlation between frequency of offending and medical comorbidity severity,³³ to our knowledge this association has never been explored in terms of HIV.

New strategies are urgently needed to sustain community treatment and align public

safety with public health goals. To support the development of gender-informed HIV interventions among CJ populations and to evaluate the specific effect of sex on HIV treatment and CJ outcomes, we analyzed data from a retrospective cohort of HIV-infected prisoners and jail detainees using longitudinally linked demographic, CJ, pharmacy, and laboratory data. The generated cohort is, to our knowledge, the largest of its kind in an integrated health system, and the only cohort to combine demographic and health data with specific data on criminal offense charges.

METHODS

The study sample was derived from the Connecticut Department of Correction (CTDOC), with approximately 16 347 inmates in 16 facilities.³⁴ Women are confined to a single facility for pretrial detainees and sentenced inmates. With the exception of 5 other states (Rhode Island, Vermont, Delaware, Alaska, Hawaii),³⁵ Connecticut has one of the few fully integrated correctional health systems that includes jails and prisons; all HIV care is provided onsite by infectious disease specialists and dedicated nurse case managers, and all Food and Drug Administration–approved ARTs are available. Although care for women is delivered in a single facility, health care across the CTDOC is standardized by the integrated health system.

The cohort has previously been described in terms of HIV treatment outcomes and recidivism.^{14,16} Individuals were included in this analysis if they (1) were incarcerated in any CTDOC facility (prison or jail) for any duration between March 2005 and June 2012, (2) had documented HIV and received ART during any incarceration, and (3) had at least 1 measurement of HIV viral load during incarceration.

Data Sources

Data were derived from 4 existing statewide sources (Figure 1): (1) a custodial database with individual demographic information and dates and types of movements into, out of, and between facilities; (2) a pharmacy database with dates and types of prescribed medications, including ART; (3) a laboratory database with CD4 and viral load measurements during incarceration; and (4) CJ information, including dates and types of all state-level criminal

offense charges. After merging databases using inmate number, we removed all unique personal identifiers to protect participant anonymity; data were stored and analyzed on triple-password-protected computers.

Measures

The primary outcome was viral suppression on entry, defined as having an HIV viral load of less than 400 copies per milliliter on intake into a CTDOC facility, to reflect HIV treatment delivered in community settings immediately preceding incarceration. Sociodemographic characteristics, derived from intake evaluations, included those listed in Table 1. We used custody information to define an incarceration period as the time from entry into a CTDOC facility to first date of release (completion of time served), conditional release to probation or parole, or death. We included incarceration period in the analysis only if HIV viral load was measured at any time. Interfacility transfers were defined as mandated movements between correctional facilities, including jails, prisons, hospitals, and halfway houses. We calculated reincarceration rate as the number of incarceration periods per time under observation, defined as the time from first release to data censor.

We characterized criminal offense charges by mutually exclusive classifications and types and subtypes. Because of the organization of the CJ data, criminal offense charges were attributable to an individual, but not to a specific incarceration period. As is customary in criminology literature, offenses were aggregated by arrest date, from which we extracted the most severe charge (on an ordinal scale,

with public disorder offenses being the least severe; followed by violation of probation or parole or failure to appear; and drug- or alcohol-related, property, and person or violent offenses being the most severe). We approximated an “offending specification” as the mean number of arrests per individual by offense classification. We also generated an offending rate, defined as the number of arrests per time under observation less the time spent incarcerated in a facility.

CJ risk scores are often used to estimate actuarial criminogenic risk and predict future related CJ involvement on the basis of a number of individual demographic and support factors and history of offending.^{36–38} Standard of practice is for courts to use these scores in sentencing decisions. We adapted a salient factor estimate of CJ risk based on the Level of Services Inventory—Revised scale³⁸ score during the observation period. We calculated scores by assigning 1 point to each of the following mutually exclusive factors:

1. having 2 or more arrests,
2. having a violation of conditional terms (i.e., violation of probation or parole or failure to appear),
3. having 2 incarcerations (2 points if ≥ 3),
4. having an escape charge,
5. ever having had a drug- or alcohol-related offense,
6. ever having had 3 or more charges on a single arrest, and
7. having the total days incarcerated within the sample’s middle tertile (2 points if highest tertile).

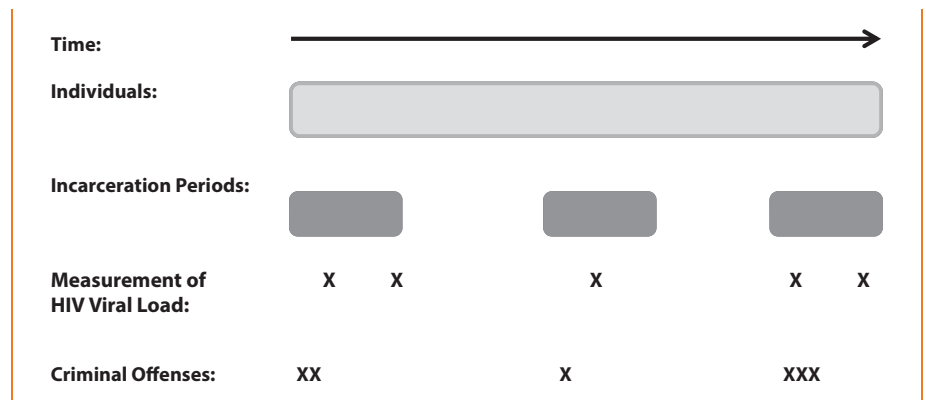


FIGURE 1—Data sources for a sample participant and levels of analysis: Connecticut Department of Corrections, 2005–2012.

TABLE 1—Descriptive Characteristics of HIV-Infected Individuals and Incarceration Periods, Stratified by Sex: Connecticut Department of Corrections, 2005–2012

Characteristics	Total Sample (n = 1089, No. Incarceration Periods = 2080)	Men (n = 866, No. Incarceration Periods = 1619)	Women (n = 223, No. Incarceration Periods = 461)	P ^a
Age, y, mean ±SD	42.6 ±8.4	43.3 ±8.5	40.2 ±7.1	< .001
Race/ethnicity, no. (%)				< .001
Non-Hispanic White	218 (20.0)	149 (17.1)	69 (31.0)	
Non-Hispanic Black	515 (47.3)	411 (47.5)	104 (46.6)	
Hispanic	351 (32.2)	303 (35.0)	48 (21.5)	
Other	5 (0.5)	3 (0.4)	2 (0.9)	
Married, no. (%)	179 (16.4)	149 (17.2)	30 (13.5)	.18
Dependent children, no. (%)				.21
None	400 (36.7)	310 (35.8)	90 (40.4)	
≥ 1	689 (63.3)	556 (64.2)	133 (59.6)	
Education, no. (%)				.28
≤ high school	502 (46.1)	392 (45.3)	110 (49.3)	
> high school	587 (53.9)	474 (54.7)	113 (50.7)	
Ever had medical insurance on entry, no. (%)	194 (17.8)	79 (9.1)	115 (51.6)	< .001
No. of incarceration periods/person, mean ±SD	1.9 ±1.4	1.9 ±1.4	2.1 ±1.4	.06
Incarceration duration, d				< .001
Mean ±SD	330.1 ±473.4	368.1 ±510.6	196.8 ±269.7	
Median (IQR)	166.5 (304)	182 (383)	96 (222)	
Interfacility transfers, no. (%)				.05
0	1954 (93.9)	1512 (93.4)	442 (95.9)	
≥ 1	126 (6.1)	107 (6.6)	19 (4.1)	
Time spent in community between incarceration periods, d ^b				
Mean ±SD	325.9 ±357.7	323.1 ±353.3	334.6 ±317.9	.3
Median (min, max)	194.5 (1, 2069)	190.5 (1, 2069)	208 (5, 2064)	.68
Intake year, no. (%)				< .001
2005–2007	1025 (49.3)	759 (46.9)	266 (57.7)	
2008–2010	803 (38.6)	655 (40.5)	148 (32.1)	
2011–2012	252 (12.1)	205 (12.7)	47 (10.2)	
Discharge status, no. (%)				.22
Probation or parole	679 (32.6)	533 (32.9)	146 (31.7)	
Release	1384 (66.6)	1070 (66.1)	314 (68.1)	
Death	17 (0.8)	16 (1.0)	1 (0.2)	
Reincarceration rate ^c				.006
Mean ±SD	0.63 ±0.92	0.67 ±1.0	0.47 ±0.38	
Median (IQR)	0.41 (0.50)	0.42 (0.52)	0.38 (0.40)	

Note. IQR = interquartile range. The sample size was n = 1089 individuals and n = 2080 incarceration periods.

^aCalculated using Student *t* test for continuous variables and χ^2 test for categorical variables unless otherwise noted.

^bFor individuals with ≥ 1 incarceration during the observation period.

^cBy Wilcoxon rank sum.

For individuals without any available offense information (26% of sampled men and 27% of sampled women), we assigned 0 points for each offense-related factor because we assumed that minor offenses would be more likely to be missing from a correctional database designed to manage the custodial population. Points were

summed, and a sample distribution histogram defined the lowest score range as 0 to 1, the middle as 2 to 3, and the highest as more than 3. When individuals with missing offense data were alternately removed from the analysis, the sample distribution of the CJ risk score and modeling results remained relatively unchanged.

We determined HIV treatment outcomes and characteristics from laboratory and pharmacy data. CD4 and viral load variables were analyzed continuously on intake and release; viral suppression was analyzed as a dichotomous variable. ART regimen was examined at the times of entry and release and was defined

by its components in terms of mutually exclusive categories, as previously described.^{14,16} We further characterized ART regimen in terms of calculated daily pill burden (based on included components and dosing strategy) and mode of medication administration, as previously described.¹⁴ We also examined whether individuals were ever prescribed ART as directly observed therapy during the observation period.

Medical and psychiatric severity scores are generated on intake by nursing staff as part of standard custodial procedure and classified on a scale ranging from 1, indicating no issues, to 5, indicating 24-hour nursing needs in a specialized unit. We grouped documented prescription of psychiatric and other medications by indicated condition and coded each category dichotomously. Diagnoses or diagnostic codes were not available in the electronic records, limiting available comorbidity data to conditions that were identified and treated, based on pharmacy records. A comorbidity score was generated for each individual participant, representing the sum total number of comorbidities (other than HIV) for which each participant was ever prescribed medication during any observed incarceration period, as shown in Table 2. Categories were mutually exclusive, and we analyzed the comorbidity score as a continuous variable.

Statistical Analysis

A sex-stratified descriptive analysis characterized individuals, incarceration periods, criminal offense charges, and treatment characteristics and outcomes, using the Student *t* test or the Wilcoxon rank-sum test to compare continuous variables and the χ^2 test for categorical variables. We modeled entry viral suppression using logistic regression with random effects (generalized linear mixed models) to account for clustered data among individuals. Models were stacked: model 1 included only key demographic characteristics (age, race/ethnicity, and education); model 2 included demographics and the CJ risk score; model 3 included demographics, the CJ risk score, and key information on medical and psychiatric comorbidities (entry ART regimen and comorbidity score); in model 4, all variables for which $P < .1$ in the bivariate analyses were entered into a multivariable model stepwise to achieve the most parsimonious

model; and model 5 included only key modifiable characteristics. We compared model fit on Akaike and Bayesian information criteria and generated separate models for men and for women. We performed all analyses using SAS version 9.3 (SAS Institute, Cary, NC).

RESULTS

The final sample of 1089 HIV-infected prisoners and jail detainees with 2080 incarceration periods included 866 (79.5%) men with 1619 incarcerations and 223 (20.5%)

TABLE 2—Characteristics of Criminal Offense Charges, Stratified by Sex: Connecticut Department of Corrections, 2005–2012

Criminal Offense Charges	Total Sample (n = 3242)	Men (n = 2631)	Women (n = 611)	P ^a
Classification, no. (%)				< .001
Misdemeanors	1322 (40.8)	1014 (38.5)	308 (50.4)	
Felonies	1235 (38.1)	1045 (39.7)	190 (31.1)	
Not classified	685 (21.1)	572 (21.7)	113 (18.5)	
Crime against persons, no (%)	463 (14.3)	401 (15.2)	62 (10.1)	< .001
Crime against property, no (%)	981 (30.3)	794 (30.2)	187 (30.6)	
Drug- or alcohol-related crime, no. (%)	624 (19.3)	530 (20.1)	94 (15.4)	.04
Possession or purchasing	318 (51.0)	261 (49.3)	57 (60.6)	
Manufacturing or sale	176 (28.2)	150 (28.3)	26 (27.7)	
Driving while Intoxicated	130 (20.8)	119 (22.5)	11 (11.7)	
Public disorder, no. (%)	1032 (31.8)	777 (29.5)	255 (41.7)	< .001
Violation of probation or parole	327 (31.7)	261 (33.6)	66 (25.9)	
Prostitution	70 (6.8)	14 (1.8)	56 (22.0)	
Failure to appear	259 (25.1)	172 (22.1)	87 (34.1)	
Other or escape	376 (36.4)	330 (42.4)	46 (18.0)	
Other types, no (%)	142 (4.4)	129 (4.9)	13 (2.1)	
Most severe arrest				< .001
No.	2111	1717	394	
Persons, no. (%)	368 (17.4)	319 (18.6)	49 (12.4)	
Property, no. (%)	761 (36.0)	618 (36.0)	143 (36.3)	
Drug or alcohol-related, no. (%)	511 (24.2)	435 (25.3)	76 (19.3)	
Violation of conditional terms, ^b no. (%)	118 (5.6)	82 (4.8)	36 (9.1)	
Other public disorder, no. (%)	353 (16.7)	263 (15.3)	90 (22.8)	
No. of arrests per individual, by class, mean \pm SD				
Total	4.06 \pm 3.3	4.13 \pm 3.3	3.77 \pm 3.1	.21
Persons	0.58 \pm 1.0	0.63 \pm 1.1	0.38 \pm 0.8	.001
Property	1.23 \pm 1.95	1.25 \pm 2.0	1.15 \pm 1.9	.59
Drug or alcohol-related	0.78 \pm 1.2	0.83 \pm 1.2	0.58 \pm 0.9	.003
Public disorder	1.47 \pm 1.7	1.42 \pm 1.7	1.65 \pm 1.6	.12
Arrest frequency per year				.006
Mean \pm SD	0.45 \pm 0.4	0.47 \pm 0.4	0.38 \pm 0.3	
Median	0.31	0.32	0.27	
CJ risk score, no.	1089	866	223	.03
Lowest range, no. (%)	257 (23.6)	189 (21.8)	68 (30.5)	
Middle range, no. (%)	400 (36.7)	329 (38.0)	71 (31.8)	
Highest range, no. (%)	432 (39.7)	348 (40.2)	84 (37.7)	

Note. CJ = criminal justice.

^aCalculated using Student *t* test for continuous variables and χ^2 test for categorical variables unless otherwise noted.

^bViolation of conditional terms includes violation of probation or parole and failure to appear.

women with 461 incarcerations. As shown in Table 1, compared with men, women were significantly more likely to be younger and to have medical insurance on entry; a greater proportion of women than men were non-Hispanic White. Overall, women's incarcerations were shorter than men's, and they were less likely to be reincarcerated, but we found no significant sex differences in terms of number of incarceration periods per person, inter-facility transfers, time spent in the community, or discharge status.

Consistent with national data,³⁹ criminal offense charges differed significantly by sex (Table 3). Compared with 2631 offense charges levied against men, the 611 charges against women were significantly less frequently for violent person crimes and more often for public disorder offenses, the majority of which involved prostitution or technical violation of probation or parole. Although men and women were nearly equally likely to have drug- or alcohol-related offenses, women's charges more often involved drug possession or purchase with intent to use. The arrest rate for women was also significantly lower than that for men.

Main sex-stratified HIV treatment outcomes are depicted in Table 2. Although men and women were equally likely to have viral suppression on entry, women were significantly more likely to achieve viral suppression pre-release, despite insignificant differences between sexes in terms of entry ART regimen or ART daily pill burden (data not shown). A minority of ART regimens were switched during incarceration (data not shown). When examined by incarceration period, men more often had ART continuously self-administered, but men and women were equally likely to have ever been prescribed ART as directly observed therapy during any observed incarceration period. Women had higher mean intake psychiatric severity scores and were 1.7 times as likely as men to ever have been prescribed antidepressant medications during incarceration. Overall, women had a significantly greater number of non-HIV treated comorbidities than men, the majority of which were mood disorders. When analyzed as continuous variables, comorbidity and CJ risk scores had no significant direct correlation for either men or women (data not shown).

We modeled entry viral suppression for 1532 incarceration periods involving 824 unique men (Table 4). In the best-fit model by stepwise selection (model 4), the adjusted odds of having viral suppression on entry were positively correlated with older age, increasing comorbidity score, and more contemporary intakes. Black and Hispanic men were significantly less likely than White men to have viral suppression on entry, and individuals with the highest calculated CJ risk were half as likely as those with the lowest CJ risk score to have viral suppression on entry.

We modeled entry viral suppression separately for 454 incarceration periods involving 219 unique women (Table 5). In the best-fit model (model 1) that included only demographic characteristics, each increasing year of age was associated with a 10% increased odds of having viral suppression on entry. Otherwise, among women, viral suppression was not directly associated with our measure of CJ risk, ART regimen, or medical or psychiatric comorbidity. When only modifiable characteristics were included as possible covariates in model 5, women's entry viral suppression was directly associated with increasing comorbidity score and shorter incarceration duration.

DISCUSSION

In this retrospective longitudinal cohort of 1089 prisoners and jail detainees with HIV on ART with 2080 incarceration periods spanning more than 7 years, we identified key differences between the sampled men and women in terms of major CJ characteristics and HIV treatment outcomes. Compared with men, women were detained for significantly shorter periods and had lower calculated criminogenic risk. Men and women entered the CTDOC with similar levels of viral suppression, reflective of their (suboptimal) community HIV treatment, but a higher proportion of women achieved viral suppression during incarceration, despite women having a higher burden of medical and psychiatric comorbidities. Findings illuminate the need for gender-informed HIV treatment strategies that can achieve maximal effectiveness among CJ-involved PLWHA.

By incorporating, for the first time to our knowledge, an in-depth examination of criminal offense charges and HIV-related health

outcomes, we uncovered some striking contrasts between men and women. Specifically, women were significantly more likely than their male counterparts to be charged with public disorder offenses, the majority of which were for prostitution or violation of conditional terms. These lower level charges might serve as a targeted opportunity to effectively engage these women in communities and better align safety goals related to correctional supervision (i.e., probation or parole) with public health goals related to retention in HIV care and ART adherence. This specific finding would otherwise have been missed in most analyses that rely more broadly on incarceration data to reflect the extent of CJ involvement.

Overall, compared with national data from the Bureau of Justice Statistics, violent crimes (including murder, rape and sexual assault, robbery, and assault) were markedly lower in this cohort of PLWHA (15% vs 54% for men and 10% vs 37% for women).³⁹ As has been shown elsewhere,⁴⁰ PLWHA are more likely to be incarcerated for minor offenses, especially those that can be more effectively addressed in the community.⁴¹

Community-based strategies have the additional benefit of reducing public spending because incarceration, especially of patients with multiple medical and psychiatric comorbidities, is expensive, and conditions are more cost effectively addressed in community settings.⁴² Community-based alternatives to incarceration strategies, which include drug and mental health courts, have key public policy implications for curbing the United States' astronomical incarceration rate. Existing evidence has suggested that drug court participation significantly reduces relapse to substance use and rates of criminal behavior and durably and cost-effectively improves overall quality of life.⁴³ Little to no data are available, however, on the effects of drug courts on health outcomes, leading some to suggest that drug courts lack transparency or are not sufficiently health centered.⁴⁴ Rather than exclude individuals with co-occurring disorders, which is common practice,⁴⁵ drug courts will be maximally effective when they align public safety and public health by incorporating treatment of other medical and social comorbidities such as HIV, psychiatric disorders, and homelessness. This shift will likely preferentially affect women

TABLE 3—Treatment and Outcomes of HIV and Other Comorbidities During Incarceration, Stratified by Sex: Connecticut Department of Corrections, 2005–2012

Characteristics	Total, ^a No. (%) or Mean ±SD	Men, ^b No. (%) or Mean ±SD	Women, ^c No. (%) or Mean ±SD	P
HIV treatment and outcomes				
HIV viral suppression ^d				
On entry	650 (32.7)	516 (33.7)	134 (29.5)	.1
On release	769 (70.6)	621 (68.7)	148 (80.0)	.002
CD4 count				
On entry	344.5 ±234.6	341.9 ±235.2	353.4 ±237.0	.36
On release	449.6 ±272.6	448.3 ±277.5	456.0 ±247.8	.73
No. of individuals ever prescribed DOT	582 ±53.4	454 ±52.4	128 ±57.4	.18
Type of medication administration ^e				<.001
Never DOT	794 (72.9)	682 (75.4)	112 (60.5)	
Transitional DOT	18 (1.7)	16 (1.8)	2 (1.1)	
Continuous DOT	25 (2.3)	15 (1.7)	10 (5.4)	
Transitional SAT	252 (23.1)	191 (21.1)	61 (33.0)	
Entry ART regimen				
PI-based	970 (48.8)	750 (49.0)	220 (48.5)	.85
nNRTI-based	730 (36.8)	554 (36.2)	176 (38.8)	.31
FDC	240 (12.1)	196 (12.8)	44 (9.7)	.08
NRTI- based	197 (9.9)	155 (10.1)	42 (9.3)	.59
INSTI	59 (3.0)	50 (3.3)	9 (2.0)	.16
Other	30 (1.5)	23 (1.5)	7 (1.5)	.95
Medical and psychiatric comorbidities				
Intake medical severity	3.4 ±0.8	3.4 ±0.8	3.4 ±0.7	.77
Intake psychiatric severity	2.8 ±1.3	2.8 ±1.3	3.1 ±0.9	
Individuals ever prescribed				
Antipsychotic	216 (19.8)	166 (19.2)	50 (22.4)	.28
Antidepressant	446 (41.0)	311 (35.9)	135 (60.5)	<.001
Both antipsychotic and antidepressant	164 (15.1)	123 (14.2)	41 (18.4)	.12
Individuals ever prescribed medications related to:				
Hypertension	371 (34.1)	314 (36.3)	57 (25.6)	.003
Diabetes	126 (11.6)	114 (13.2)	14 (6.3)	.004
Dyslipidemia	117 (10.7)	103 (11.9)	14 (6.3)	.02
Seizure disorder	178 (16.4)	144 (16.6)	34 (15.3)	.62
Herpes simplex virus	213 (19.6)	156 (18.0)	57 (25.6)	.01
Asthma	229 (21.0)	146 (16.9)	83 (37.2)	<.001
Neuropathy	44 (4.0)	25 (2.9)	19 (8.5)	<.001
HCV	20 (1.8)	18 (2.1)	2 (0.9)	.4
Opioid dependence	94 (8.6)	68 (7.9)	26 (11.7)	.07
Pregnancy	13 (1.2)	...	13 (5.8)	...
Comorbidity score	1.9 ±1.7	1.8 ±1.6	2.3 ±1.6	<.001

Note. ART = antiretroviral therapy; DOT = directly observed therapy; FDC = fixed-dose combination; INSTI = integrase strand transfer inhibitor; nNRTI = non-nucleoside reverse transcriptase inhibitor; NRTI = nucleoside reverse transcriptase inhibitor; PI = protease inhibitor; SAT = self-administered therapy.

^aThe sample size was n = 1089 individuals and 2080 incarceration periods.

^bThe sample size was n = 866 men and 1619 incarceration periods.

^cThe sample size was n = 223 women and 461 incarceration periods.

^dNot all incarceration periods had viral loads available for both entry and release, so the percentage is the number with viral suppression/the number with available viral load on entry or on release, respectively.

^eNot all pharmacy prescriptions specified whether the medication was provided as DOT or SAT, so the denominator is based on the available information.

TABLE 4—Correlates of Viral Suppression on Entry Among Men: Connecticut Department of Corrections, 2005–2012

Covariate	Model 1, AOR (95% CI)	Model 2, AOR (95% CI)	Model 3, AOR (95% CI)	Model 4, AOR (95% CI)
Age (continuous)	1.06* (1.04, 1.09)	1.06* (1.04, 1.09)	1.06* (1.03, 1.08)	1.05* (1.02, 1.07)
Race/ethnicity				
Non-Hispanic White (Ref)	1.00	1.00	1.00	1.00
Non-Hispanic Black	0.31* (0.18, 0.52)	0.30* (0.18, 0.51)	0.32* (0.19, 0.53)	0.35* (0.21, 0.58)
Hispanic	0.35* (0.20, 0.61)	0.35* (0.20, 0.60)	0.35* (0.21, 0.61)	0.37* (0.22, 0.62)
Education				
≤ high school (Ref)	1.00	1.00	1.00	...
> high school	0.84 (0.57, 1.23)	0.83 (0.37, 1.03)	0.83 (0.57, 1.21)	...
CJ risk score				
Lowest range (Ref)	...	1.00	1.00	1.00
Middle range	...	0.82 (0.48, 1.41)	0.75 (0.44, 1.27)	0.81 (0.47, 1.39)
Highest range	...	0.62 (0.37, 1.03)	0.55* (0.33, 0.92)	0.49* (0.29, 0.82)
Entry ART regimen				
PI based	0.35 (0.11, 1.13)	...
nNRTI based	0.47 (0.14, 1.57)	...
FDC	0.59 (0.34, 1.00)	0.63 (0.38, 1.03)
NRTI only	0.34 (0.10, 1.18)	...
INSTI	1.27 (0.30, 5.43)	2.11 (0.87, 5.14)
Comorbidity score	1.11* (1.00, 1.24)	1.13* (1.01, 1.26)
Incarceration period by intake year				
2005–2007 (Ref)	1.00
2008–2010	2.13* (1.52, 2.97)
2011–2012	2.83* (1.74, 4.58)
Medical insurance on entry				
Yes (Ref)	1.00
No	0.56 (0.31, 1.03)
AIC/BIC	1815.79/1844.06	1815.77/1853.46	1806.90/1872.86	1788.45/1835.57

Note. AOR = adjusted odds ratio; AIC = Akaike information criterion; ART = antiretroviral therapy; BIC = Bayesian information criterion; CI = confidence interval; CJ = criminal justice; DOT = directly observed therapy; FDC = fixed-dose combination; INSTI = integrase strand transfer inhibitor; nNRTI = nonnucleoside reverse transcriptase inhibitor; NRTI = nucleoside reverse transcriptase inhibitor; PI = protease inhibitor; SAT = self-administered therapy. Model 1 includes demographic characteristics only (age, race/ethnicity, education); model 2 includes demographics and CJ risk; model 3 includes demographics, CJ risk, and HIV and other comorbidity severity; model 4 includes all covariates for which $P < .1$ on bivariate association, followed by stepwise selection. The sample size was $n = 824$ individuals and $n = 1532$ incarceration periods. * $P < .05$.

who, as shown here, experience a significantly higher burden of comorbidity than men. Instead of using drug courts to off-load prisons, alternatives to incarceration strategies can be leveraged to help stabilize chaotic lives in a more effective and enduring way than imprisonment, in which HIV treatment outcomes are temporarily optimized, then deteriorate rapidly after release.^{14,16}

Previous published literature from the criminology and legal fields has suggested that

individuals with untreated medical comorbidities are more likely to commit crimes and ultimately become CJ involved. In a study of 915 offenders in Canada, for example, diabetes was twice as prevalent as in the general population, was mostly undiagnosed or untreated at the time of offense, and was most frequent among violent and sex offenders whose victims were children.³³ From a legal perspective, uncontrolled diabetes has historically been considered a disease of the mind and within the

scope of the insanity defense,⁴⁶ although the association between medical comorbidity severity and criminality or aggression has more recently been suggested in relation to epilepsy,⁴⁷ dyslipidemia,⁴⁸ and other diseases, mostly in men. Missing from these studies has been consideration of race/ethnicity, neighborhood, and socioeconomic status as confounders that likely contribute both to serious health disparities and to CJ risk.⁴⁹ We found, for example, that Black and Hispanic men were significantly less likely than their White counterparts to have viral suppression on entry. Even after we controlled for these potential confounders in multivariable models, however, men with the highest levels of CJ risk were consistently and least likely to have viral suppression.

Our findings suggest that the association, though not causative, between CJ risk and poor health outcomes extends to include HIV. Unlike diabetes or epilepsy, HIV is a chronic disease that is generally asymptomatic. HIV differs from other chronic diseases in a more important way because when it is inadequately treated and viral suppression is not achieved, the virus can be transmitted to others in the setting through risk behaviors.⁵⁰ We identify here a population of PLWHA who are disengaged from major systems of health care and social support, putting them at higher risk of CJ involvement and suboptimal HIV treatment outcomes. Criminogenic risk and HIV viral suppression are each modifiable characteristics amenable to targeted interventions. Correctional community supervision—that is, probation or parole—may provide an opportunity for intervention from both public health and public safety perspectives. Costs and criminal behaviors decrease when people can be successfully maintained in the community.

For women in this cohort, the array of available data was insufficient to fully explain viral suppression on entry using various analytical methods. Although the smaller sample of women may have underpowered our analyses, we suspect instead that a number of unmeasured confounders may have biased results toward the null. Women's pathways to substance use, criminality, and incarceration, unlike those of their male counterparts, are more often related to lack of social or economic capital.^{51–54} Thus, for women, CJ risk is most strongly associated with comorbid conditions

TABLE 5—Correlates of Viral Suppression on Entry Among Women: Connecticut Department of Corrections, 2005–2012

Covariate	Model 1, AOR (95% CI)	Model 2, AOR (95% CI)	Model 3, AOR (95% CI)	Model 4, AOR (95% CI)	Model 5, AOR (95% CI)
Age (continuous)	1.10* (1.04, 1.17)	1.10* (1.04, 1.17)	1.09* (1.03, 1.16)	1.09* (1.03, 1.15)	...
Race/ethnicity					
Non-Hispanic White (Ref)	1.00	1.00	1.00
Non-Hispanic Black	1.17 (0.50, 2.71)	1.15 (0.49, 2.69)	1.18 (0.35, 3.28)
Hispanic	1.11 (0.37, 3.30)	1.11 (0.37, 3.31)	1.08 (0.50, 2.81)
Education					
≤ high school (Ref)
> high school	0.75 (0.35, 1.60)	0.96 (0.43, 2.14)	0.79 (0.37, 1.69)
CJ risk score					
Lowest range (Ref)	...	1.00	1.00	...	1.00
Middle range	...	0.68 (0.25, 1.89)	0.60 (0.21, 1.71)	...	0.80 (0.30, 2.12)
Highest range	...	0.71 (0.29, 1.75)	0.59 (0.22, 1.56)	...	0.57 (0.23, 1.42)
Entry ART regimen					
PI-based	0.56 (0.06, 4.88)
nNRTI-based	0.35 (0.04, 3.29)
FDC					
NRTI only	0.43 (0.04, 4.40)
INSTI	1.02 (0.05, 19.21)
Comorbidity score	1.13 (0.89, 1.43)	...	1.24* (0.98, 1.56)
Incarceration period by intake year					
2005–2007 (Ref)	1.00	...
2008–2010	1.67 (0.90, 3.09)	...
2011–2012	2.03 (0.77, 5.35)	...
Medical insurance at entry					
Yes (Ref)	1.00
No	0.72 (0.27, 1.91)
Time incarcerated, mo	0.96* (0.92, 1.00)
Discharge status					
Conditional release (Ref)	1.00
Release/death	0.97 (0.48, 1.94)
Ever prescribed DOT					
No (Ref)	1.00
Yes	0.81 (0.39, 1.68)
AIC/BIC	516.39/536.67	519.67/546.70	527.26/574.58	516.69/543.73	537.90/568.40

Note. AIC = Akaike information criterion; AOR = adjusted odds ratio; ART = antiretroviral therapy; BIC = Bayesian information criterion; CI = confidence interval; CJ = criminal justice; DOT = directly observed therapy; FDC = fixed-dose combination; INSTI = integrase strand transfer inhibitor; nNRTI = nonnucleoside reverse transcriptase inhibitor; NRTI = nucleoside reverse transcriptase inhibitor; PI = protease inhibitor. Model 1 includes demographic characteristics only (age, race/ethnicity, education); model 2 includes demographics and CJ risk; model 3 includes demographics, CJ risk, and HIV and other comorbidity severity; model 4 includes all covariates for which $P < .1$ on bivariate association, followed by stepwise selection; model 5 includes only modifiable characteristics. The sample size was $n = 219$ individuals and $n = 454$ incarceration periods.

* $P < .05$.

and involvement in sex work as well as related partner dependency, partner violence,²⁶ and homelessness—factors that were not well represented or measured in the databases we used.

Compared with CJ-involved men, CJ-involved women have a higher prevalence of unrecognized or untreated psychiatric and substance use disorders,⁵⁵ which may contribute to their

criminal behavior. To disentangle this association, we first have to understand the extent to which untreated comorbidities in the community contribute to the relatively low proportion of women with viral suppression on entry into the CJ setting,¹⁹ and develop detailed assessments during incarceration and after release.

We have previously reported that women

released from jail were significantly more likely than men to report ongoing cocaine use and were half as likely as men to sustain viral suppression 6 months after release.¹⁹

Given the synergistic effect of substance use, violence, psychiatric disorders, and social instability on HIV-related health outcomes of CJ-involved women, sex-specific and culturally

relevant interventions will incorporate a syndemic approach that collectively addresses social and medical conditions.^{23,56} Although several previous syndemic HIV prevention interventions have been described,^{57–59} lack of sustainability remains a major concern.⁶⁰ To maximize immediate and durable benefits for health and CJ outcomes, evidence-based interventions could be adapted and incorporated into existing structural systems, such as probation and parole.⁶¹

Limitations

Although this retrospective cohort study provides important new insight into multidimensional and sex-specific factors affecting prisoner health, analyses were necessarily restricted by available data. In some cases, unvalidated scales were used, either because they are routinely used by correctional systems to manage custodial populations and determine service needs (e.g., intake medical or psychiatric severity scores) or because they were calculated composite scales (e.g., comorbidity score, CJ risk score) that required imputations for missing data. Although we used all available data to sufficiently represent offense history, we could not account for federal offense charges or charges acquired out of state that may have affected outcomes, though we suspect this represented a minority of the sample; community-based data on health outcomes or service utilization were also not available. Our findings likely overestimate the proportion of inmates with HIV who achieve viral suppression because data were restricted only to those diagnosed and prescribed ART. Finally, generalizability might be limited to this state's correctional health system, though we suspect, given the large sample size and long period of observation, that critical issues raised here are likely relevant to other systems of CJ and health care delivery.

Conclusions

To our knowledge, from the largest contemporary cohort of HIV-infected inmates on ART, women's incarceration and HIV treatment outcomes differed in critical and dynamic ways from men's, providing insight into addressing sex-related health disparities. Women's higher prevalence of nonviolent offenses and increased medical and psychiatric comorbidities, especially treatable substance

use disorders, supports alternatives to incarceration strategies to holistically provide treatment. Interventions for HIV-infected CJ populations should be sex specific to effectively align health and justice goals. ■

About the Authors

Jaimie P. Meyer and Frederick L. Altice are with the AIDS Program, Yale School of Medicine, New Haven, CT. Jaimie P. Meyer is also with the Chronic Disease Epidemiology Department, Yale School of Public Health, New Haven. Javier Cepeda and Frederick L. Altice are with the Department of Epidemiology of Microbial Diseases, Yale School of Public Health. Faye S. Taxman is with the Criminology, Law, and Society Department, George Mason University, Fairfax, VA.

Correspondence should be sent to Jaimie P. Meyer, 135 College Street, Suite 323, New Haven, CT 06510 (e-mail: jaimie.meyer@yale.edu). Reprints can be ordered at <http://www.ajph.org> by clicking the "Reprints" link.

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Contributors

J. P. Meyer had full access to all of the data in the study and was responsible for the integrity of the data and the accuracy of the data analysis. J. P. Meyer and F. L. Altice provided funding support and generated the study concept and design, with input from F. S. Taxman. J. P. Meyer and J. Cepeda acquired the data, and J. Cepeda performed the analysis; all authors interpreted the data. J. P. Meyer drafted the article, with critical revision for important intellectual content by J. Cepeda, F. S. Taxman, and F. L. Altice. All authors saw and approved the final version for publication.

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Human Participant Protection

All procedures were independently approved by Yale's institutional review board and the Connecticut Department of Corrections Research Advisory Committee.

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