

# Immunity to HAV and/or HBV Among Inmates Living with HIV

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## BACKGROUND

- Hepatitis A (HAV) and B viruses (HBV) are vaccine-preventable diseases.
- Incarcerated patients have greater risk factors for acquiring both HAV and HBV, such as intravenous drug use, high risk sexual activity, and tattoos.<sup>1</sup>
- The prevalence of HBV is **five times greater** in those who are incarcerated than in the general population, and there are no guidelines mandating routine HBV vaccinations for inmates.<sup>2,3</sup>
- Recent studies have shown that providing vaccinations is a more effective and less costly approach to decrease the burden of these diseases.<sup>4</sup>
- Screening upon entry to prison provides an ideal public health opportunity to assess vaccination status and administer vaccination while incarcerated.

## RESEARCH DESIGN AND METHODS

- Retrospective, electronic medical record review of incarcerated adults receiving human immunodeficiency virus (HIV) telemedicine care in 26 prisons in Illinois, USA, between January 1, 2019 through December 31, 2019.
- Statistical analysis included Chi-squared testing and descriptive statistics.

### Inclusion Criteria:

- Adults ≥18 years of age
- Confirmed diagnosis of symptomatic or asymptomatic HIV/AIDS
- Incarcerated in the Illinois Department of Corrections (IDOC) during the retrospective study period
- Available data for HAV/HBV serologies, HIV-1 viral load, and CD4 count during incarceration

### Exclusion Criteria:

- Age <18 years
- No documented HAV/HBV serologies, HIV-1 viral load, and CD4 count during incarceration

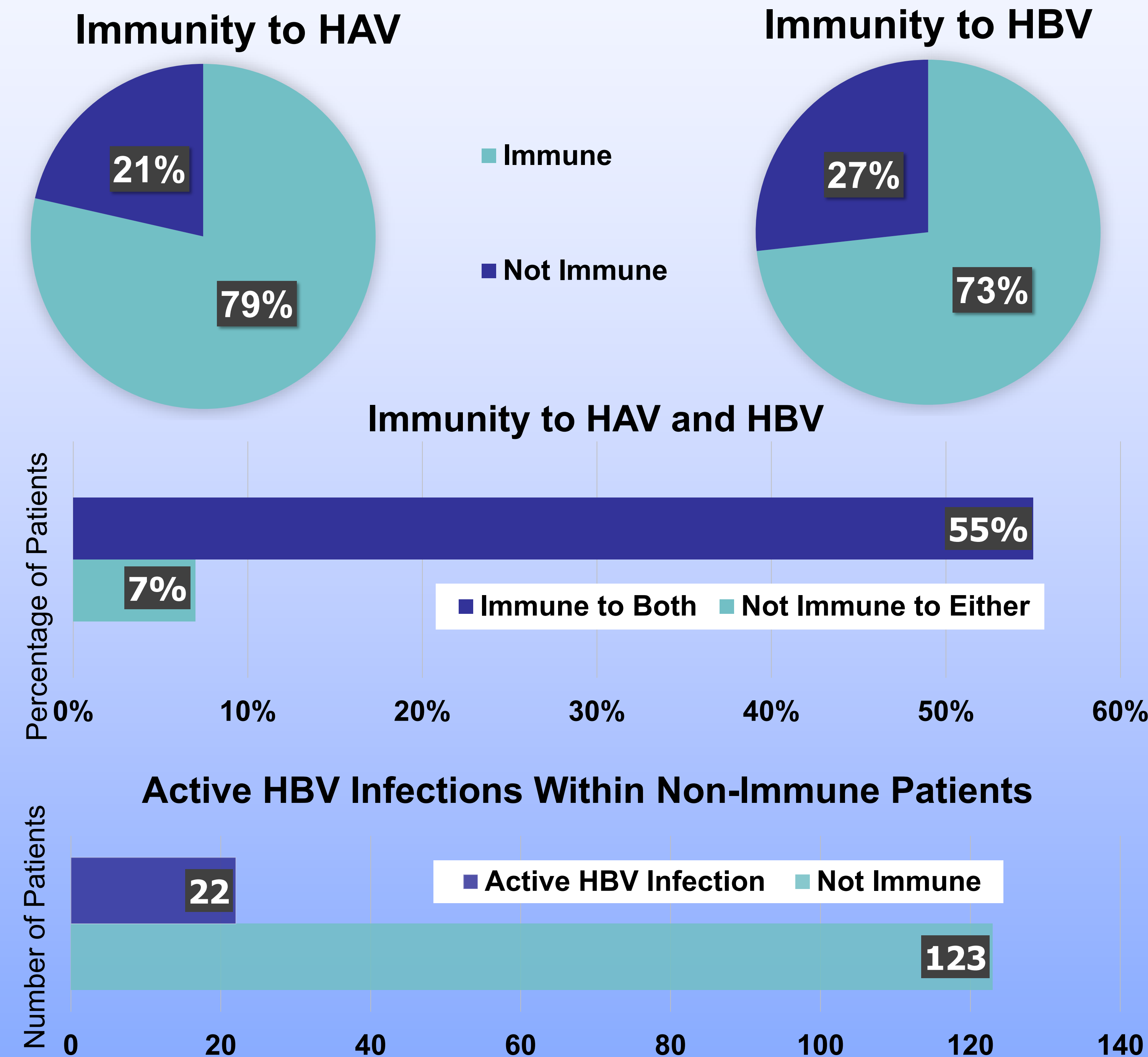
## ENDPOINTS

- Primary Objective:** assess rates of HAV and/or HBV immunity in individuals living with HIV in IDOC.
- Secondary Objective:** assess factors associated with vaccination status.

## BASELINE DEMOGRAPHICS

Characteristic	Category	Number of Patients
Age	18-50	348 (67.1%)
	51-100	171 (32.9%)
Sex	Male	480 (92.5%)
	Female	39 (7.5%)
Race	Black	392 (75.5%)
	White	72 (13.9%)
	Latino	29 (5.6%)
	Other	26 (5.0%)
CD4 Count	≤200 cells/mm <sup>3</sup>	26 (5.0%)
	>200 cells/mm <sup>3</sup>	493 (95%)
CD4%	0-25%	153 (29.5%)
	26-50%	350 (67.4%)
	51-75%	16 (3.1%)
Viral Load	Detectable (>20 copies/mL)	78 (15%)
	Undetectable (≤20 copies/mL)	441 (85%)

## RESULTS



- Among the 524 patients analyzed, the majority were Black men (75%) with an average age of 44 years.
- 397 patients had existing data for HBV vaccination, where 5% had HBV infection, 1.4% had an equivocal HBV surface antibody and negative HBV surface antigen, and 70% had documented immunity.
- In patients who had a detectable HIV-1 viral load, 16 were not immune to HAV, 17 were not immune to HBV, and 3 were not immune to both HAV and HBV.
- In patients who had a CD4 count of >200 cells/mm<sup>3</sup>, 88 were not immune to HAV, 117 were not immune to HBV, and 26 were not immune to both HAV and HBV.
- Immunity did not vary based on CD4 count, age, gender, or race (p > 0.05).**

## CONCLUSIONS

- Providing HAV and HBV vaccinations to the incarcerated can increase immunity and reduce transmission within this vulnerable population.
- This is of particular importance for patients living with HIV as this is an indication for vaccination.
- Based on these findings, the telemedicine study team has been able to assess serologies and advocate for vaccination for inmates living with HIV entering the IDOC.
- Over time, we expect our interventions to result in further improvements in rates of immunity.

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## DISCLOSURES

The authors of this presentation have no disclosures concerning possible financial or personal relationships with commercial entities that may have a direct or indirect interest in the subject matter of this presentation.