

#### Comparison of weight changes in treatment-naïve HIV-infected patients receiving **1701 N Capitol Ave** integrase inhibitor-based therapy compared to protease inhibitor-based therapy **Indianapolis, IN 46202** Zachary Howe, PharmD, Eric Farmer, PharmD, BCPS, AAHIVP, Emily Huesgen, PharmD, BCACP, AAHIVP, **Contact:** zhowe@iuhealth.org

## Background

Current guidelines recommend integrase inhibitor-ba (INSTI) therapy as first line in treatment-naïve patien with HIV. However, recent data indicate they may be associated with increases in weight, BMI, and body Protease inhibitors (PI) are known to alter metabolism body weight and are a potential alternative regimen is certain clinical situations.<sup>5-8</sup>

Variation in clinical outcomes related to weight gain been observed with differing demographic factors<sup>9</sup> and nucleoside reverse transcriptase inhibitor (NRTI) backbones utilized<sup>10</sup> alongside INSTIs. Published dat bictegravir-based regimens are scarce, and data comp these drug classes could inform clinical decision mak based on baseline patient characteristics.

# Methods

- Retrospective observational cohort
- Data collected from a chart review for all treatment naïve patients initiating a PI- or INSTI-based regin from 1/1/13 to 7/31/19.
- Exclusion criteria: Less than 18 years old, less than months of therapy, pregnancy, quadriplegia or paraplegia, amputation, or lack of weight or lab dat 10-14 months after starting ART.



## **Statistical Analysis**

Categorical Variables: Chi Square Analysis with or without Bonferroni correction (\*), as appropriate

**Continuous Variables:** Wilcoxon Rank Sum Test

**Power:** In order to detect a difference of 7%, assuming a baseline average weight of  $76 \pm 15 \text{ kg}^4$  with a power of 80% and type 1 error rate of 5%, 282 patients were needed.

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	INSTI	PI		INSTI	D
	(N=145)	(N=17)		(N=145)	(N=1
Sex			<b>INSTI or PI</b>		
Male	82.8%	76.5%	ABC/3TC	26.2%	5.99
Female	17.2%	23.5%	NDF	37.2%	88.2
			TAF	36.6%	5.99
Ethnicity			p value	0.0034*	
Asian American	0.7%	11.8%	ATV	10.00/	17.6
Hispanic	2 1%	5.9%	BIC	10.3%	00.4
	<i>2</i> .170	5.770		26 60/	82.4
African American	57%	47%		30.0% 40.70/	
Caucasian	40%	35.3%		49.7%	
			KAL	3.4%	
			Results		
		Me	edian Weight Gair	1	
6			(p-0.32)	<b>7</b> 1	
5		3 (		5.1	
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	r 10	Protease II		inically Cignifican	t Waisht
Median Weight Gain (n=0.021*)			U	Gain (p=0.03	*)
8	7 7	· /	60.0%	50.0% 53.6%	
Sun 6	5	.1	40.0%		
250 10 10		2.4	20.0% -		23.7%
2					
		$\Delta RC/2TC$	0.0%	TAF TDF A	BC/3TC
	$\mathbf{M} = 1 \mathbf{D} \mathbf{\Gamma}$	BN	II Category Chang	ges	
1000/			(p=0.0008*)	100%	
100%					



![](_page_0_Figure_18.jpeg)

In conjunction with the findings of previous studies, it would seem some amount of weight gain with INSTI- or PI-based therapy can be expected for some patients. Additionally, baseline characteristics, such as sex, race, comorbidities, and background therapies may accentuate or attenuate this effect.

Contrary to other studies, both forms of tenofovir were associated with increased weight gain and a higher incidence of clinically significant weight gain, rather than only being associated with TAF. This study considered the potential impact of baseline characteristics on incidence and degree of weight gain and provided data for newer agents, such as bictegravir. The small number of patients receiving PI-based therapy and dyssynchrony between prescribing practices during the study time period and current practice limit the applicability of these findings.

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#### Discussion

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