

Characteristics of Patients Living with HIV (PLWH) and HIV Treatment Outcomes in Community Practices in Southern vs Non-Southern US Regions

¹Keith Rawlings, MD, ²Paul Sax, MD, ¹Julie Priest MSPH, ³Janna Radtchenko, MBA, ¹Joe Mrus, MD, ⁴Joe Eron, MD, ¹Alan Oglesby, MPH, ⁵Keri Althoff, PhD, ⁶Faith Fletcher, PhD, ⁷Moti Rampogal, MD, ⁸Steven Santiago, MD, ⁹Richard A Elion, MD

¹ViiV Healthcare, Research Triangle, NC, USA, ²Brigham and Women's Hospital and Harvard Medical School, Boston, MA, USA, ³Trio Health Analytics, La Jolla, CA, USA, ⁴University of North Carolina at Chapel Hill, Chapel Hill, NC, USA, ⁵John Hopkins University Bloomberg School of Public Health, Baltimore, MD, USA, ⁶University of Alabama at Birmingham School of Public Health, Birmingham, AL, USA, ⁷Midway Specialty Care Center, Fort Pierce, FL, USA, ⁸Care Resource, Miami Beach, FL, USA, ⁹George Washington University, Washington, DC, USA



1. BACKGROUND

This study evaluated individual characteristics and antiretroviral therapy (ART) prescribing for people with HIV (PWH) seeking care at practices in southern vs non-southern US regions.

2. METHODS

PWH (>18 years old) starting a new ART between January 2015-September 2019 with viral load at first regimen prescription (baseline), and ≥6 months of prior ART history (if not treatment-naïve), were selected from Trio Health HIV EMR database.

Comparisons of baseline characteristics were conducted via chi-square for categorical variables and t-test for continuous variables.

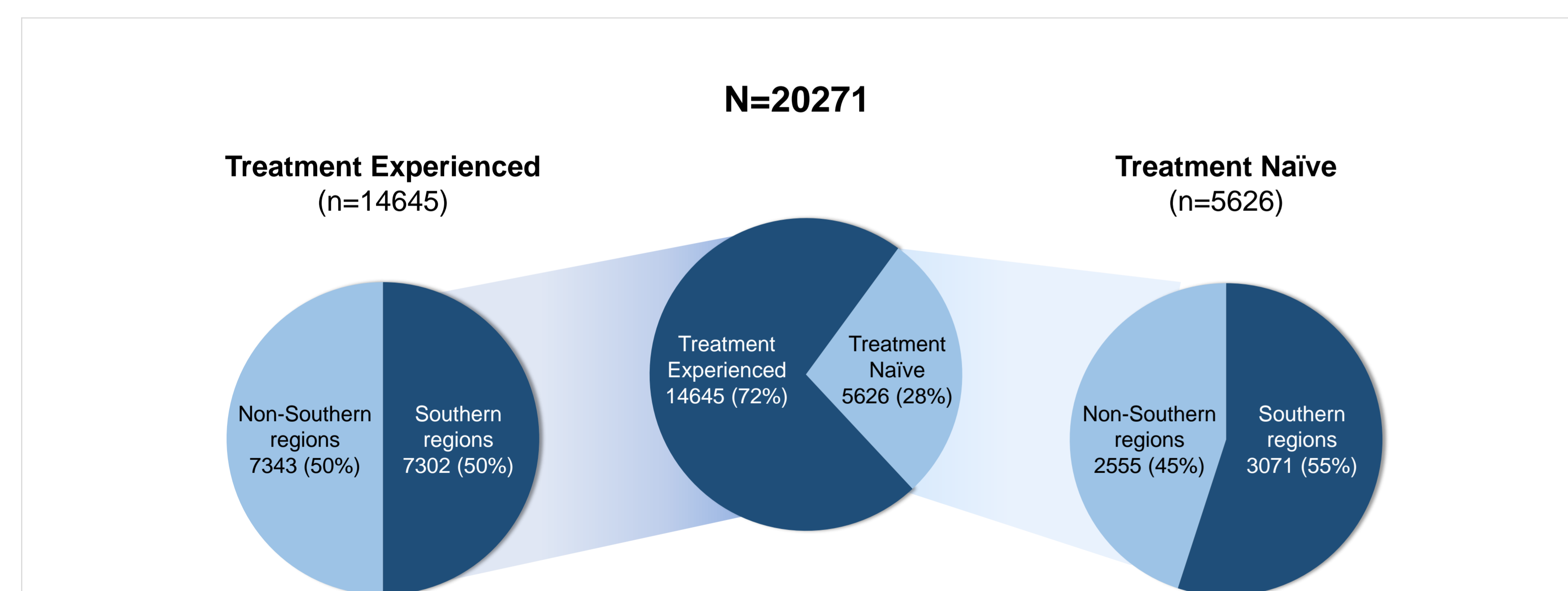
Multivariate logistic regression with binary outcome "viremic at last observation" estimated the associations of covariates among those with viral load recorded ≥12 months after baseline (n=12,689).

Regions were defined per US Census and sample availability (South included: TX, FL; Non-South: IL, NM, CA, PA).

3. RESULTS

Of 20,271 PWH, 14,645 (72%) were treatment-experienced, 7,302, 50% in South) and 5,626 (28%) treatment-naïve (3,071, 55% in South) [Figure 1].

FIGURE 1. PATIENT SELECTION



n (%) unless specified	Treatment Naïve (N=5626)			Treatment Experienced (N=14645)			
	Southern (n=3071)	Non-Southern (n=2555)	p-value	Southern (n=7302)	Non-Southern (n=7343)	p-value	
Age, mean (SD)	39.1 (12)	37.4 (11.9)	<0.001	47.4 (11.9)	46.6 (12.1)	<0.001	
Age >50	651 (21)	448 (18)	0.001	3160 (43)	3091 (42)	0.148	
Male	2290 (75)	2005 (78)	<0.001	5748 (79)	5877 (80)	<0.001	
Race	White	1461 (48)	1035 (41)	<0.001	3934 (54)	3946 (54)	<0.001
	Black	1218 (40)	990 (39)		2256 (31)	1920 (26)	
	Other	223 (7)	140 (5)		493 (7)	402 (5)	
	Unknown	169 (6)	390 (15)		619 (8)	1075 (15)	
Payer Type	Commercial	1014 (33)	1618 (63)	<0.001	3543 (49)	5296 (72)	<0.001
	Medicare	171 (6)	186 (7)		1177 (16)	988 (13)	
	Medicaid	121 (4)	243 (10)		306 (4)	397 (5)	
	Other	648 (21)	477 (19)		899 (12)	622 (8)	
	Unknown	1117 (36)	31 (1)		1377 (19)	40 (1)	

This study was supported by ViiV Healthcare

Baseline characteristics of treatment-naïve and treatment-experienced groups differed by region [Table 1-2].

Among treatment-experienced, 83% were suppressed at baseline in South vs. 91% in non-South (p<0.001); after ≥12 months, suppression rates were 85% vs. 91%, p<0.001, respectively.

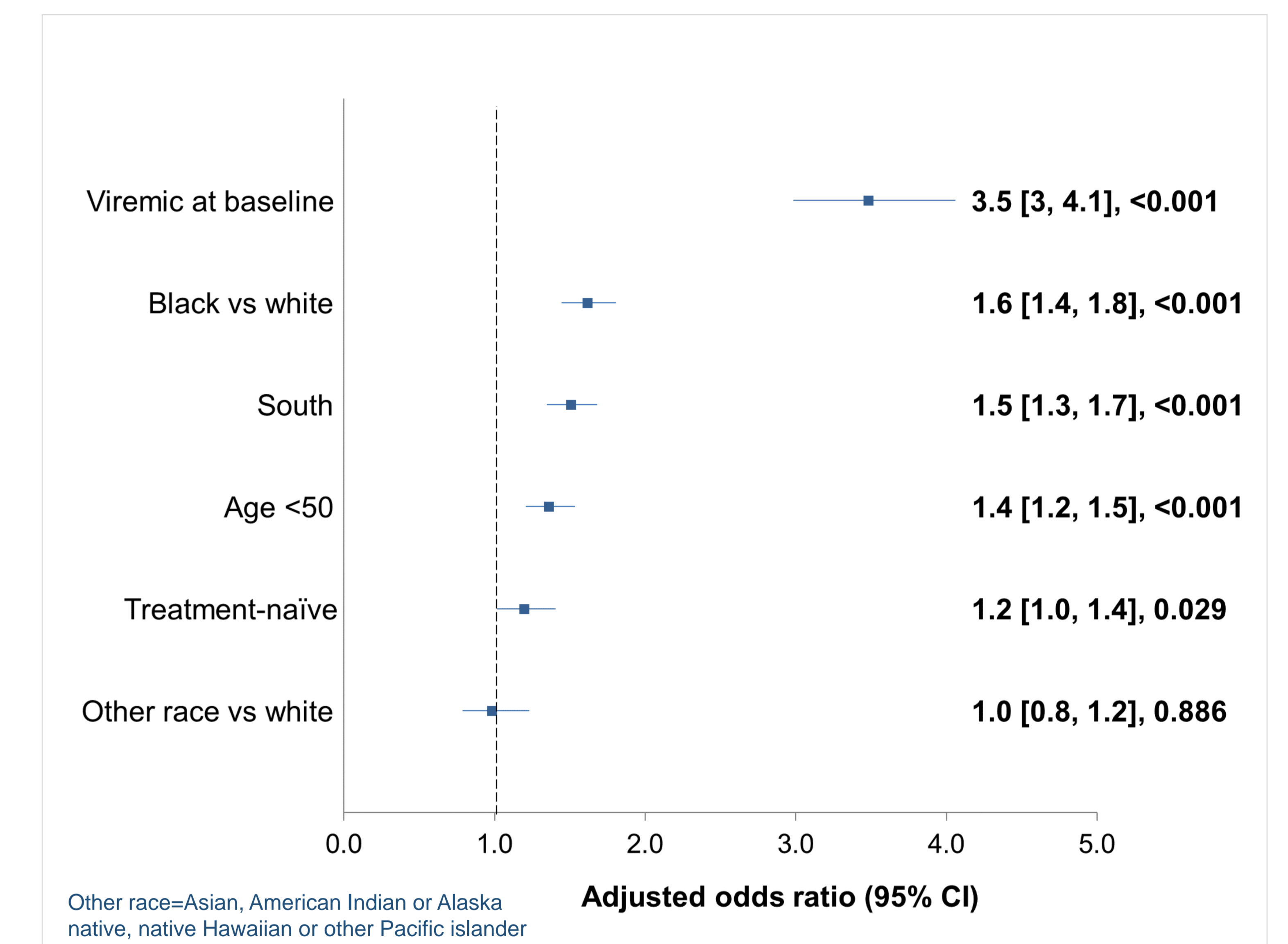
Among treatment-naïve with ≥12 months of follow-up, 72% were suppressed in South, 76% in non-South (p=0.008).

Treatment-experienced individuals in the South were less likely to be on a single-tablet regimen than non-South (60% vs. 62%, p=0.021), no differences by region among treatment-naïve (59% vs 58%, p=0.470).

n (%) unless specified	Treatment Naïve (N=5626)			Treatment Experienced (N=14645)			
	Southern (n=3071)	Non-Southern (n=2555)	p-value	Southern (n=7302)	Non-Southern (n=7343)	p-value	
Charlson Comorbidity Index, mean (SD)	3.5 (3.2) n=3071	4.3 (2.9) n=2555	<0.001	5.3 (2.9) n=7302	5.5 (3) n=7343	<0.001	
Hemoglobin, mean (SD)	13.8 (2) n=2484	13.9 (1.9) n=1639	0.571	14.4 (1.7) n=6055	14.6 (1.7) n=5799	<0.001	
Platelets, mean (SD)	131.8 (119.5) n=2528	205.9 (101.9) n=1513	<0.001	147.5 (116.8) n=6026	224.5 (77.5) n=5771	<0.001	
Protein Urine, mean (SD)	10.1 (17) n=932	11.2 (17.7) n=783	0.200	7.9 (11.5) n=3409	11.4 (20.9) n=2392	<0.001	
BMI	Underweight	132 (5) n=2697	84 (4) n=2394	<0.001	181 (3) n=6574	127 (2) n=7017	<0.001
	Normal	1189 (44) n=2697	1156 (48) n=2394		2099 (32) n=6574	2665 (38) n=7017	
	Overweight	825 (31) n=2697	758 (32) n=2394		2475 (38) n=6574	2680 (38) n=7017	
	Obese	551 (20) n=2697	396 (17) n=2394		1819 (28) n=6574	1545 (22) n=7017	
CD4 <200 cells/ml	488 (22) n=2177	308 (27) n=1161	0.008	343 (6) n=5885	262 (5) n=4854	0.335	
eGFR	<60	82 (3) n=2516	79 (3) n=2483	0.004	636 (10) n=6278	629 (9) n=7222	<0.001
	60-89	592 (24) n=2516	489 (20) n=2483		2676 (43) n=6278	2759 (38) n=7222	
	90+	1842 (73) n=2516	1915 (77) n=2483		2966 (47) n=6278	3834 (53) n=7222	
FIB-4	<1.3	1052 (43) n=2444	967 (64) n=1519	<0.001	2645 (44) n=6009	3558 (65) n=5463	<0.001
	1.3-3.25	308 (13) n=2444	278 (18) n=1519		1028 (17) n=6009	1472 (27) n=5463	
	>3.25	1084 (44) n=2444	274 (18) n=1519		2336 (39) n=6009	433 (8) n=5463	
	Hyperglycemia (glucose ≥1.1 g/l)	93 (11) n=875	200 (16) n=1277		0.001	278 (18) n=1508	
Triglycerides ≥1.5 g/l	561 (33) n=1721	334 (29) n=1135	0.074	1802 (42) n=4338	1814 (39) n=4693	0.005	
Hypertension (≥130 mmHg systolic, ≥90 mmHg diastolic)	297 (12) n=2494	185 (10) n=1775	0.131	778 (12) n=6342	403 (10) n=4167	<0.001	
HDL-cholesterol <40 mg/dl males, <50 mg/dl females	727 (48) n=1509	753 (48) n=1561	0.973	1611 (38) n=4289	1762 (35) n=5099	0.003	
Arteriosclerotic Cardiovascular Disease (ASCVD) risk >7.5%	96 (20) n=486	74 (31) n=235	0.001	877 (36) n=2461	645 (46) n=1406	<0.001	
Suppressed at baseline	0 (0)	0 (0)	N/A	6083 (83)	6686 (91)	<0.001	
Cardiovascular Disease (CVD)	534 (17)	1077 (42)	<0.001	2330 (32)	3411 (46)	<0.001	
Diabetes	64 (2)	91 (4)	0.001	542 (7)	618 (8)	0.026	
Hypertension	331 (11)	281 (11)	0.792	1672 (23)	2035 (28)	<0.001	
Renal	40 (1)	36 (1)	0.730	511 (7)	457 (6)	0.059	

In the logistic regression accounting for race, gender, age, practice region, treatment status, and baseline viral suppression, individuals younger than 50, black or African American vs. white, treated in South region, treatment-naïve, or viremic at baseline were more likely to be viremic after ≥12 months. Final model is shown in Figure 2.

FIGURE 2. PREDICTORS OF VIREMIA AT 12 MONTHS



4. LIMITATIONS

Limitations of this study are typical of retrospective observational studies: subjects were non-randomized, observers were non-blinded, and some subgroups are small in sample size.

Viral suppression rates at 6 and 12 months since baseline were evaluated regardless of regimen.

Data is limited to treatment centers captured in the Trio database and may not represent treatment patterns and patient characteristics in the entire US. All patients were treated at nationally qualified health centers.

5. CONCLUSION

These data suggest population differences and treatment disparities between Southern and other US regions. Policy makers should examine structural and societal reasons for these disparities in an effort to reduce or eliminate them.

M. Keith Rawlings, Joe Mrus, Julie Priest, and Alan Oglesby are employed by ViiV Healthcare. Paul E. Sax consults for Gilead Sciences, ViiV Healthcare, Merck, Janssen. He received research grants from Gilead Sciences, Merck, and ViiV Healthcare. Janna Radtchenko is employed by Trio Health. Joseph J. Eron consults for Merck, ViiV Healthcare, Gilead Sciences, and Janssen. The University of North Carolina receives research funding from ViiV Healthcare, Gilead Sciences, and Janssen from which he receives support as an investigator. Keri N. Althoff previously served on a Medical Advisory Board for Gilead Sciences. Moti Rampogal is a consultant for ViiV Healthcare, Gilead Sciences, Janssen, and Merck and received research funding from these companies. He is a speaker for ViiV Healthcare, Gilead Sciences, and Janssen. Steven Santiago serves on the Medical Advisory Board for Gilead and is a Speaker for Gilead and Janssen. Richard A. Elion received grants from Gilead Sciences and Proteus, serves on the Advisory boards for Gilead Sciences and ViiV Healthcare, and is a speaker for Gilead Sciences and Janssen. Drs. Althoff, Eron, Eron, Santiago, and Sax serve on Trio Health's Scientific Advisory Board.