Table S2: Characteristics of treatment-experienced patients receiving a second or subsequent ART regimen (n=2866)

Variables		ΡΙ	INSTI
Variables	NNRTI (n=716)	(n=1575)	(n=575)
Age at inclusion (in years)	(H=710)	(11–1373)	(11-373)
Mean (SD)	43.5 (9.8)	43.0 (9.7)	44.3 (11.5)
Sex (%)	43.3 (9.8)	43.0 (9.7)	44.3 (11.3)
Male	613 (85.6%)	1264 (80.2%)	495 (86.1%)
Risk factor for HIV acquisition	013 (63.070)	1204 (80.278)	493 (80.170)
MSM (%)	449 (62.7%)	949 (60.2%)	399 (69.4%)
Bisexual (%)	18 (2.5%)	51 (3.2%)	11 (1.9%)
Heterosexual (%)	154 (21.5%)	380 (24.1%)	82 (14.3%)
From endemic countries (%)	113 (15.8%)	241 (15.3%)	56 (9.7%)
Vertical transmission (%)	9 (1.3%)	10 (0.6%)	4 (0.7%)
Delay in ART treatment initiation (in	7 (212.13)	20 (01012)	(**, ***)
years)			
Mean (SD)	1.7 (3.4)	1.8 (3.6)	1.7 (3.9)
ART duration before inclusion (in years)	` /	` ,	, ,
Mean (SD)	5.8 (4.7)	6.3 (5.3)	5.8 (5.9)
Time since diagnosis at inclusion (in			
years)			
Mean (SD)	7.2 (6.0)	7.9 (6.3)	7.2 (7.1)
Treatment changes before inclusion			
Yes	332 (46.4%)	873 (55.4%)	174 (30.3%)
Viral load (copies/ml) at inclusion			
<50	277 (38.7%)	475 (30.2%)	181 (31.5%)
50-10000	209 (29.2%)	496 (31.5%)	168 (29.2%)
>10000	175 (24.4%)	520 (33.0%)	204 (35.5%)
MD	55 (7.7%)	84 (5.3%)	22 (3.8%)
Median (25%-75%)	270.0 (49.5-12082.0)	462.0 (49.5-38000.0)	601.0 (49.5-38400.0)
CD4 count at inclusion (cells/mm3)	00 (10 10 ()		
<200	89 (12.4%)	362 (23.0%)	70 (12.2%)
200-350	207 (28.9%)	440 (27.9%)	149 (25.9%)
>350	362 (50.6%)	684 (43.4%)	327 (56.9%)
MD	58 (8.1%)	89 (5.7%)	29 (5.0%)
Median (25%-75%)	381.5 (260.0-580.0)	330.0 (207.0-500.0)	410.0 (279.0-597.0)
CD8 count at inclusion (cells/mm3)	250 (26 20/)	615 (39.0%)	202 (35.1%)
<800	259 (36.2%) 312 (43.6%)	727 (46.2%)	202 (33.1%) 235 (40.9%)
≥800	,	` /	
MD Median (25%-75%)	145 (20.2%) 850.0 (600.0-1150.0)	233 (14.8%) 850.0 (582.0-1210.0)	138 (24.0%) 841.0 (610.0-1180.0)
Ratio CD4/CD8 at inclusion	830.0 (000.0-1130.0)	830.0 (382.0-1210.0)	841.0 (610.0-1180.0)
<1	498 (69.6%)	1259 (79.9%)	387 (67.3%)
>1	73 (10.2%)	82 (5.2%)	49 (8.5%)
MD	145 (20.2%)	234 (14.9%)	139 (24.2%)
Median (25%-75%)	0.4 (0.2-0.6)	0.4 (0.3-0.7)	0.5 (0.3-0.7)
CD4 nadir before inclusion	0.1 (0.2 0.0)	0.1 (0.3 0.7)	0.5 (0.5 0.7)
<200	124 (17.3%)	457 (29.0%)	99 (17.2%)
200-350	235 (32.8%)	490 (31.1%)	168 (29.2%)
>350	299 (41.8%)	539 (34.2%)	279 (48.5%)
MD	58 (8.1%)	89 (5.7%)	29 (5.1%)
Median (25%-75%)	337.5 (230.0-510.0)	281.0 (180.0-427.0)	360.0 (240.0-500.0)
Hepatitis B before inclusion	(== 3.0 5 20.0)		(= 1010 0000)
Positive for HBsAg	45 (6.3%)	80 (5.1%)	16 (2.8%)
Negative for HBsAg	541 (75.6%)	1189 (75.5%)	464 (80.7%)
Not documented	130 (18.1%)	306 (19.4%)	95 (16.5%)
Hepatitis C before inclusion	(()	
Positive for anti-HCV	46 (6.4%)	190 (12.1%)	29 (5.0%)
Negative for anti-HCV	374 (52.2%)	753 (47.8%)	343 (59.7%)
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Not documented	296 (41.4%)	632 (40.1%)	203 (35.3%)
Cytomegalovirus serostatus			
Positive	172 (24.0%)	387 (24.6%)	110 (19.1%)
Negative	32 (4.5%)	82 (5.2%)	49 (8.5%)
Not documented	512 (71.5%)	1106 (70.2%)	416 (72.4%)
Previous exposure to mono/dual NRTI therapy before inclusion			
Yes	53 (7.4%)	291 (18.5%)	57 (9.9%)
Previously documented virologic failure			
before inclusion			
Yes	21 (2.9%)	166 (10.5%)	21 (3.7%)
Year of inclusion			
2006-2009	517 (72.2%)	1268 (80.5%)	161 (28.0%)
2010-2013	176 (24.6%)	286 (18.2%)	226 (39.3%)
2014-2017	23 (3.2%)	21 (1.3%)	188 (32.7%)

Abbreviations: NNRTI, non-nucleoside reverse-transcriptase inhibitor; NRTI, nucleoside reverse-transcriptase inhibitor; NRTI, nucleoside reverse-transcriptase inhibitor; NRTI, nucleoside reverse-transcriptase inhibitor; NRTI, Integrase Strand Transfer Inhibitor; ART: Antiretroviral therapy; SD, standard deviation; HIV, human immunodeficiency virus; MSM, men who have sex with men; MD, missing data; HBsAg, hepatitis B surface antigen; HVC, hepatitis C virus;