		Expt/ culture #	P# (Days <sup>c</sup> )	Gag region <sup>a,b</sup>					
Start Virus <sup>a</sup>	Start [drug], xEC50			Capsid				Capsid/SP1	
				218	219	326	332	362	364
V362I/ V370A	1	3/1,2.3	P11		H219Q (2/3)				
V362I	1	3/1,2.3	P11		H219Q		T332T/N (1/3)		
T332S/ V362I	1	3/1,2.3	P11		H219Q (2/3)				
T332S	1	3/1	P11		H219Q			V362V/I	
		3/2			H219Q				A364V
		3/3			H219P				A364V
R286K/ V370A	1	3/1	P11		H219Q			V362I	
		3/2		V218V/A		A326T		V362I	
		3/3				A326T		V362I	
R286K	1	3/2	P11		H219Q			V362V/I	A364A/V
		3/1,3			H219Q				A364V
∆V370	1	3/1,2.3	P11		H219Q	A326A/T			
ΔT371	1	3/1,2.3	P11		H219Q				A364A/V

S1 Table: Amino Acid Substitutions Selected in Gag During Independent Sequential Passage Experiments in the Presence of Increasing Concentrations of GSK35323795

<sup>a</sup>WT virus was NL<sub>4-3</sub>RepRluc Gag P373S, variant viruses were NL<sub>4-3</sub>RepRluc Gag P373S with the indicated additional changes in Gag. Infections were initiated at a multiplicity of infection of 0.005; <sup>b</sup> sequence changes were observed in Matrix, SP2/P6 or protease genes; <sup>c</sup>Number of days in culture. EC<sub>50</sub>, 50% effective concentration; P#, passage number; SP1, spacer peptide 1; WT, wild-type