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Determining the Frequency and Mechanisms of HIV-1 and HIV-2 RNA Copackaging by Single-Virion Analysis[∇]

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HIV-1 and HIV-2 are derived from two distinct primate viruses and share only limited sequence identity. Despite this, HIV-1 and HIV-2 Gag polyproteins can coassemble into the same particle and their genomes can undergo recombination, albeit at an extremely low frequency, implying that HIV-1 and HIV-2 RNA can be copackaged into the same particle. To determine the frequency of HIV-1 and HIV-2 RNA copackaging and to dissect the mechanisms that allow the heterologous RNA copackaging, we directly visualized the RNA content of each particle by using RNA-binding proteins tagged with fluorescent proteins to label the viral genomes. We found that when HIV-1 and HIV-2 RNA are present in viral particles at similar ratios, ~10% of the viral particles encapsidate both HIV-1 and HIV-2 RNAs. Furthermore, heterologous RNA copackaging can be promoted by mutating the 6-nucleotide (6-nt) dimer initiation signal (DIS) to discourage RNA homodimerization or to encourage RNA heterodimerization, indicating that HIV-1 and HIV-2 RNA can heterodimerize prior to packaging using the DIS sequences. We also observed that the coassembly of HIV-1 and HIV-2 Gag proteins is not required for the heterologous RNA copackaging; HIV-1 Gag proteins are capable of mediating HIV-1 and HIV-2 RNA copackaging. These results define the *cis*- and *trans*-acting elements required for and affecting the heterologous RNA copackaging, a prerequisite for the generation of chimeric viruses by recombination, and also shed light on the mechanisms of RNA-Gag recognition essential for RNA encapsidation.

The two etiological agents of AIDS, human immunodeficiency virus type 1 (HIV-1) and HIV-2, were introduced into the human population by zoonotic transmission of simian immunodeficiency viruses (SIVs) that naturally infect chimpanzees or gorillas (SIV_{cpz} or SIV_{gor}) and sooty mangabeys (SIV_{sm}), respectively (12, 13, 17, 19, 26, 30, 32). Consequently, these lentiviruses, which are the only two known to infect humans, share only limited sequence identity (~48% nucleotide [nt] identity at the genome level and ~60% amino acid identity between their Gag polyproteins). Although both viruses can cause AIDS, HIV-1 is more pathogenic in humans and is distributed worldwide, whereas HIV-2 infection is limited geographically and mostly located in West Africa (20). However, infection with one of the HIVs does not provide protection from infection with the other HIV. Indeed, there is a significant population of people who are dually infected with both HIV-1 and HIV-2 (11, 14). HIV-1 and HIV-2 target the same cell types, using both the same receptor and the same coreceptors for virus entry (1, 31, 34). The possible interplay between these two viruses has been an intriguing topic of research aimed at understanding the extent and the mechanisms of such molecular interactions.

One unique feature of retrovirus replication is that two copies of the full-length RNA are packaged into a viral particle. The specific packaging of the viral RNA, which makes up less than 1% of the total cellular RNA, is mediated by at least two determinants. One determinant is the viral structural protein Gag and especially the nucleocapsid domain in the polypro-

tein. The second determinant is the packaging signal in the viral RNA, most of which resides at the 5' end of the viral genome, including part of the 5' untranslated region, and often extends into part of *gag*. The interactions between Gag and viral RNA ensure the selection and encapsidation of the proper RNAs into the particles. Retroviral RNA packaging is generally specific and limited to the RNAs of the same virus or closely related viruses. However, cross-packaging does occur, including two known examples of nonreciprocal RNA packaging of distantly related viruses; one set is HIV-1 and HIV-2 (18), and the other set is murine leukemia virus (MLV) and spleen necrosis virus (SNV) (3). SNV Gag polyprotein can efficiently package both SNV and MLV RNA; however, MLV Gag polyprotein does not package SNV RNA. Similarly, it has been shown that HIV-1 Gag can package HIV-2 RNA, whereas HIV-2 Gag does not package HIV-1 RNA (18). However, certain aspects of RNA packaging mechanisms used by HIV-1 and HIV-2 are similar. HIV-1 RNA has been shown to dimerize or select its copackaged RNA partner prior to encapsidation (4, 23, 24). A palindromic sequence located at the top of stem-loop 1 (SL1), termed the dimerization initiation signal (DIS), plays an important role during the RNA partner selection process (6, 8). HIV-1 RNAs that have complementary DIS sequences can copackage together more efficiently than those that do not (7, 23, 28). It is thought that the DIS sequences of the two RNAs form intermolecular base pairings to initiate the dimerization or the RNA partner selection process. Our recent studies showed that the RNA packaging mechanisms of HIV-2 have many features similar to those of HIV-1, including RNA partner selection prior to encapsidation and the involvement of a palindromic sequence in this process (27).

It has been shown that HIV-1 and HIV-2 can interact molecularly in several ways. For example, despite sharing a low

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level of amino acid sequence identity, the Gag polyproteins from HIV-1 and HIV-2 can coassemble into the same particle and complement each other's functions to carry out infection (2). It has also been shown in a cell culture system that HIV-1 and HIV-2 can recombine and generate hybrid genomes, albeit at a rate far lower than that occurring between two HIV-1 or between two HIV-2 genomes (25). Retroviral recombination is the result of the reverse transcriptase (RT) switching between the two copackaged RNA templates during DNA synthesis. Therefore, the observation that HIV-1 and HIV-2 can recombine indicates that the RNA genomes of these two different viruses must be able to be packaged into the same particle (heterologous RNA copackaging). However, it was unclear how frequently these heterologous RNAs can be copackaged and whether elements in the viral RNAs exist that constrain such copackaging. Furthermore, it is unclear whether the coassembly of the HIV-1 and HIV-2 Gag polyproteins is required to promote copackaging of the heterologous RNAs by each interacting with its own RNA.

In this study, we used the single-virion analysis for direct measurement of the frequency of HIV-1 and HIV-2 RNA copackaging. This system, based on labeling HIV-1 and HIV-2 genomes with fluorescent protein-tagged RNA-binding proteins, offers a direct measurement of viral RNA content at single-virion resolution. Unlike standard biochemical assays that measure the efficiency with which a given RNA is packaged, this microscopy-based method allows us to determine whether two RNAs are copackaged into the same virion. We found that when HIV-1 and HIV-2 RNAs are packaged at similar levels, ~10% of the particles contained both HIV-1 and HIV-2 RNA. The heterologous RNA copackaging frequency can be influenced by the identities of the DIS sequences, indicating that HIV-1 and HIV-2 RNAs can heterodimerize using the DIS sequences prior to packaging. We also determined that HIV-1 Gag alone is sufficient for heterologous RNA copackaging.

MATERIALS AND METHODS

Viral vectors and plasmid construction. The HIV-1 constructs GagCeFP-MS2SL and Gag-MS2SL have been previously described (4); for clarity, in this report these plasmids are referred to as HIV-1-GagCeFP-MS2SL and HIV-1-Gag-MS2SL, respectively. Briefly, these two constructs were derived from the molecular clone NL4-3; portions of the *pol*, *vif*, *vpr*, *vpu*, and *env* genes were deleted, rendering the encoded proteins nonfunctional. Additionally, 24 copies of the stem-loop recognized by the bacteriophage MS2 coat protein were inserted in the *pol* gene. All *cis*-acting elements required for HIV-1 replication, including the packaging signal, are present in these genomes. The two plasmids have similar structures, except the HIV-1-GagCeFP-MS2SL expresses Gag tagged with cerulean fluorescent protein (CeFP), whereas HIV-1-Gag-MS2SL expresses untagged Gag. In all experiments, tagged Gag-CeFP and untagged Gag are coexpressed to avoid possible distortions of viral particle morphology. For brevity, only GagCeFP-expressing constructs are mentioned.

The mutant with altered DIS sequences, HIV-1-DIS6C-GagCeFP-MS2SL, was derived from HIV-1-GagCeFP-MS2SL by changing the GCGCGC sequence in the DIS into CCCCCC. This was done by replacing the SphI-XhoI DNA fragment in 6CDIS-GagCeFP-BglSL with the corresponding fragment from HIV-1-GagCeFP-MSL; both the Gag-CeFP and the untagged Gag versions were generated (4). A Gag mutant, HIV-1-noGag-MSL, which contains a stop codon after amino acid 109 of CA, was generated by introducing a 4-bp frameshift mutation in the *gag* gene of HIV-1-Gag-MS2SL. This was achieved by digesting HIV-1-Gag-MS2SL with SpeI, followed by a fill-in reaction using the Klenow fragment of *Escherichia coli* polymerase and DNA ligation.

The HIV-2 constructs 2-GagCeFP-BglSL, 2-Gag-BglSL, and 2-6G-GagCeFP-BSL have been previously described (27); for clarity, these constructs are re-

ferred to as HIV-2-GagCeFP-BglSL, HIV-2-Gag-BglSL, and HIV-2-DIS6G-GagCeFP-BglSL in this report. These two constructs were derived from the ROD12 molecular clone and contained all *cis*-acting elements important for viral replication and inactivating mutations in *pol*, *vpr*, and *env*. Additionally, both constructs contained 18 copies of stem-loops (BglSL) in *pol* recognized by the *E. coli* BglG protein. Construct HIV-2-GagCeFP-BglSL expresses Gag tagged with CeFP, whereas HIV-2-Gag-BglSL expresses untagged Gag (27). As with HIV-1, only the GagCeFP construct is mentioned, although both CeFP-tagged HIV-2 Gag and wild-type HIV-2 Gag were coexpressed in all experiments.

Construct HIV-2-GagCeFP-BglSL-noTatRev was generated by digesting HIV-2-GagCeFP-BglSL plasmid DNA with BsmBI, followed by a fill-in reaction using the Klenow fragment from *E. coli* DNA polymerase and DNA ligation. This procedure generated an inactivating frameshift mutation in both *tat* and *rev*. The HIV-2-DIS6G-GagCeFP-BglSL-noTatRev construct was generated using the same procedure by starting with HIV-2-DIS6G-GagCeFP-BglSL plasmid. The HIV-2-noGag-BglSL was derived from a previously described 2⁺T-17 mutant that contained two stop codons in *gag*, one at codon 17 of Gag in MA and one at codon 171 of CA (27). DIS mutants in the HIV-2-noGag-BglSL context were constructed using 2-step PCR. Primers harboring the DIS mutations were used to amplify the HIV-2-noGag-BglSL template, yielding a 2.4-kb DNA fragment containing DIS mutations and both stop codons in Gag that was cloned into the HIV-2-GagCeFP-BglSL backbone by the use of the NgoMIV and SmaI restriction sites. The general structures of the newly generated plasmids were determined by restriction enzyme mapping, and regions generated by PCR were characterized by DNA sequencing to avoid inadvertent mutations.

The MS2-yellow fluorescent protein (MS2-YFP) construct was a generous gift from Robert Singer (Albert Einstein Medical College); the Bgl-mCherry plasmid has been previously described (4).

Cell culture, transfection, and virus production. The 293T human embryonic kidney cell line was grown in a humidified 37°C incubator with 5% CO₂ and maintained in Dulbecco's modified Eagle's medium supplemented with 10% fetal bovine serum, penicillin (50 U/ml), and streptomycin (50 µg/ml). In all experiments, plasmids expressing Gag and GagCeFP were cotransfected at a 1:1 molar ratio; DNA transfections were performed using FuGeneHD (Roche) according to manufacturer's recommendations. Supernatants were harvested from 293T cells 19 to 20 h posttransfection, clarified through a 0.45-µm-pore-size filter, and either stored at -80°C or used immediately for image acquisition.

Single-virion analyses. To prepare particles for microscopy, clarified culture supernatant was mixed with Polybrene and the mixture was plated on a 35-mm-diameter glass-bottom plate (MatTek Corp.) and incubated at 37°C for 2 h. An inverted Nikon Eclipse Ti microscope with a 100× oil objective and a numerical aperture of 1.40 and an X-Cite 120 system (EXFO Photonic Solutions Inc.) were used for image capture as previously described (27). The filter settings were 427/10 nm and 480/40 nm (CeFP), 504/12 nm and 542/27 nm (YFP), and 577/25 nm and 632/60 nm (mCherry) for excitation and emission, respectively. Digital images were acquired by the use of an AndOr technology iXon camera and NIS Element AR software (Nikon). Custom software developed with Matlab was used to identify particles and their colocalization with RNA.

RESULTS

Determining HIV-1 and HIV-2 RNA genome copackaging frequency. To measure the frequency of HIV-1 and HIV-2 RNA copackaging, we used two previously described constructs that express near-full-length HIV-1 and HIV-2 genomes containing stem-loop sequences recognized by RNA-binding proteins (Fig. 1A) (4, 27). These viral genomes allowed us to fluorescently tag and directly visualize encapsidated RNA in individual viral particles. The modified HIV-1 genome HIV-1-GagCeFP-MS2SL expresses HIV-1 Gag tagged with CeFP, Tat, and Rev; additionally, this construct also contains 24 copies of MS2SL, the stem-loop sequence specifically recognized by the bacteriophage MS2 coat proteins. The modified HIV-2 genome, HIV-2-GagCeFP-BglSL, expresses viral gene products, including HIV-2 Gag tagged with CeFP, Tat, and Rev; additionally, it contains 18 copies of BglSL, the stem-loop sequence specifically recognized by the *E. coli* BglG protein. Two versions of each viral construct were generated; one ex-

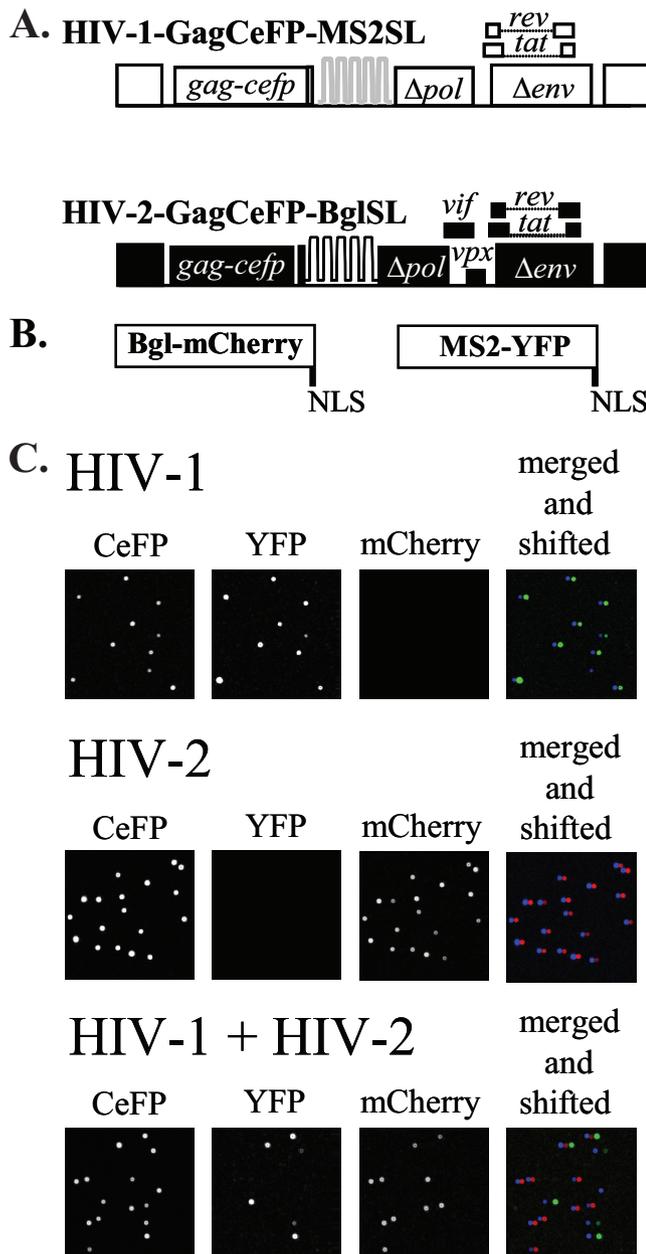


FIG. 1. System used to determine HIV-1 and HIV-2 RNA copackaging efficiency. (A) General structures of the HIV constructs. HIV-1 construct HIV-1-GagCeFP-MS2SL contains in its RNA stem-loops recognized by the bacteriophage MS2 coat protein. HIV-2 construct HIV-2-GagCeFP-BglISL contains in its RNA stem-loops recognized by the *E. coli* BglG protein. Both constructs express Gag tagged with CeFP. In all experiments, these constructs were coexpressed with sister constructs that were identical except that they expressed wild-type Gag proteins without CeFP. For simplicity, the constructs expressing wild-type Gag are not shown. White boxes represent HIV-1 sequences, whereas black boxes represent HIV-2 sequences. (B) General structures of RNA-binding proteins tagged with fluorescent proteins containing nuclear localization signals (NLS) at the C terminus. (C) Representative images of single-virion analyses. MS2-YFP and Bgl-mCherry were cotransfected along with viral constructs in all samples. All three channels were merged and shifted (the YFP and mCherry channels were shifted 5 and 10 pixels to the right, respectively) as shown in the fourth column to illustrate the RNA signals associated with each particle.

presses wild-type Gag and the other expresses Gag tagged with CeFP. In all experiments, GagCeFP and Gag were coexpressed to preserve normal particle morphology; for simplicity, only the names of the GagCeFP constructs are mentioned.

To examine the RNA content in the HIV particles, viral constructs were transfected into 293T cells along with two plasmids that express fluorescently tagged RNA-binding proteins, MS2-YFP and Bgl-mCherry (Fig. 1B). The supernatant was harvested 19 to 20 h posttransfection and clarified, and images of viral particles were obtained using fluorescence microscopy. Examples of images of viral particles obtained from 293T cells transfected with HIV-1-GagCeFP-MS2SL, MS2-YFP, and Bgl-mCherry are shown in the upper panels of Fig. 1C. As a portion of the HIV-1 Gag polyproteins were tagged with CeFP, viral particles can be identified by their signals in the CeFP channel. The full-length RNA expressed by HIV-1-GagCeFP-MS2SL contained sequences recognized by the MS2 coat protein; as a result, the majority (>90%) of the CeFP⁺ particles also contained YFP signals but not mCherry signals (summarized in Table 1). In contrast, HIV-2 construct HIV-2-GagCeFP-BglISL contains sequences recognized by Bgl proteins; hence, most of the CeFP⁺ particles generated from cotransfection of HIV-2-GagCeFP-BglISL, MS2-YFP, and Bgl-mCherry also contained mCherry signals but not YFP signals (Fig. 1C, middle panels; Table 1). These results showed that most of the HIV particles contained viral RNA genomes; furthermore, the signals observed from MS2-YFP or Bgl-mCherry were specific to the stem-loops in the viral genomes and had little background and nonspecific labeling.

We then generated particles by coexpressing HIV-1-GagCeFP-MS2SL, HIV-2-GagCeFP-BglISL, MS2-YFP, and Bgl-mCherry (Fig. 1C, lower panels). Most of the CeFP⁺ particles also had either YFP or mCherry signals, indicating that they contained either HIV-1 or HIV-2 RNA, respectively; some of the CeFP⁺ particles were positive for YFP and mCherry signals, indicating that they contained both HIV-1 and HIV-2 RNA. As the relative levels of abundance of these two viral RNAs affect the copackaging frequency, in our experiments, we used only results from experiments that showed similar levels of HIV-1 and HIV-2 RNA signals. In 13 independent experiments, between 6% and 13% of the particles in the viral population exhibited CeFP, YFP, and mCherry signals, with an average of 9.7% and standard deviation of 2.8% (Table 1).

Examining the influence of DIS on copackaging of heterologous viral RNAs. In both HIV-1 and HIV-2, RNA partner selection occurs prior to the encapsidation of the viral genome; additionally, the DIS sequences play a major role in the selection process (4, 6, 8, 23, 24, 27). However, previous experiments examined copackaging of RNAs derived from the same viral species; it is unclear whether DIS can affect copackaging of heterologous viral RNAs. We sought to determine whether RNAs from genetically distinct viruses can interact through DIS sequences and be copackaged. The sequences of the 5' untranslated regions of HIV-1 and HIV-2 have little homology; the two molecular clones we used had 42% nucleotide sequence identity, but the two sequences had similar predicted RNA structures (Fig. 2A and B). The DIS sequences for HIV-1 and HIV-2 are discordant; the subtype B HIV-1 used in this study has a GCGCGC sequence whereas HIV-2 has a

TABLE 1. Single-virion analyses of HIV-1 and HIV-2 RNA copackaging

Coexpressed constructs ^a	CeFP ⁺ particles			YFP ⁺ + mCherry ⁺ (%) ^b	RNA labeling efficiency ^c
	No. analyzed	YFP ⁺ (%)	mCherry ⁺ (%)		
HIV-1–GagCeFP-MS2SL					
Expt 1	1,005	93.2	0.0	0.0	93.2
Expt 2	1,772	93.7	0.1	0.3	94.1
Expt 3	1,786	92.9	0.0	0.1	93.0
Expt 4	2,369	92.0	0.0	0.0	92.0
Expt 5	5,032	92.8	0.0	0.0	92.8
Expt 6	5,060	91.8	0.0	0.0	91.8
Expt 7	4,568	91.6	0.0	0.0	91.6
Expt 8	5,578	91.5	0.0	0.2	91.7
Expt 9	7,505	91.3	0.0	0.3	91.6
Expt 10	3,655	92.3	0.0	0.1	92.4
Expt 11	8,138	93.0	0.0	0.0	93.0
Expt 12	20,159	91.4	0.0	0.4	91.8
Expt 13	16,632	92.0	0.0	0.0	92.0
HIV-2–GagCeFP-BglSL					
Expt 1	1,644	0.1	92.7	0.4	93.2
Expt 2	2,902	0.0	93.7	0.2	93.9
Expt 3	5,708	0.0	93.8	0.1	93.9
Expt 4	4,738	0.0	93.3	0.3	93.6
Expt 5	6,467	0.0	92.7	0.5	93.2
Expt 6	1,077	0.0	91.8	1.0	92.8
Expt 7	8,240	0.0	92.8	0.4	93.2
Expt 8	10,027	0.0	95.6	0.3	95.9
Expt 9	15,232	0.0	91.4	0.2	91.6
Expt 10	15,107	0.0	94.1	0.0	94.1
Expt 11	2,384	0.0	92.1	0.5	92.6
HIV-1–GagCeFP-MS2SL + HIV-2–GagCeFP-BglSL					
Expt 1	1,025	43.8	44.4	6.1	94.3
Expt 2	1,251	45.2	42.0	5.7	92.9
Expt 3	1,651	53.0	34.0	5.6	92.6
Expt 4	3,072	32.8	50.7	9.5	93.0
Expt 5	5,983	50.4	33.6	8.0	92.0
Expt 6	12,462	36.6	46.0	10.2	92.8
Expt 7	15,366	44.2	37.6	10.9	92.7
Expt 8	13,505	49.9	32.1	10.9	92.9
Expt 9	7,259	52.1	33.0	8.0	93.1
Expt 10	12,956	37.4	41.1	13.1	91.6
Expt 11	12,245	36.9	42.6	12.9	92.4
Expt 12	10,749	35.4	44.3	12.1	91.8
Expt 13	10,431	36.0	42.4	12.7	91.1

^a Although not indicated, both Bgl-mCherry and MS2-YFP were coexpressed in all experiments.

^b Average plus standard deviation (SD) for HIV-1-GagCeFP-MS2SL + HIV-2-GagCeFP-BglSL, 9.7 ± 2.8 .

^c Calculated by adding the values from the three columns under the CeFP⁺ heading.

GGTACC sequence (Fig. 2). Using an *in vitro* assay, it was previously shown that RNA dimerization occurs only when HIV-1 and HIV-2 RNA contain the same DIS (9). To examine whether the frequency of heterologous RNA copackaging can be altered, we generated viral constructs containing various DIS sequences and tested the abilities of their RNAs to be copackaged with RNA from another virus. We first mutated the HIV-2 DIS sequence to GCGCGC so that this mutant (HIV-2-DIS3GC-GagCeFP-BglSL; Fig. 2D) had the same DIS sequence as the HIV-1 construct HIV-1-GagCeFP-MS2SL. We found that the presence of matching and complementary DIS sequences did not have a significant effect on the frequency of heterologous RNA copackaging; approximately 12% of the particles contained both HIV-1 RNA and HIV-2

RNA with GCGCGC at the DIS (range in 7 experiments, 10% to 14%) (Table 2).

Although mutating the HIV-2 DIS sequence to GCGCGC may have allowed base pairing of the DIS sequences in the HIV-1 and HIV-2 RNA, this mutation did not interfere with the ability of two RNAs from the same virus to base pair at the DIS sequences. To simultaneously promote heterodimerization and discourage homodimerization of the viral RNAs, we mutated the HIV-1 DIS sequence to CCCCCC (HIV-1-DIS6C-GagCeFP-MS2SL; Fig. 2C) and the HIV-2 DIS sequence to GGGGGG (HIV-2-DIS6G-GagCeFP-BglSL; Fig. 2D), coexpressed these constructs with RNA-binding proteins, and examined the resulting viruses. Indeed, the copackaging frequency of these two RNAs increased to ~31% (range in 9

TABLE 2. The effects of DIS sequences on HIV-1 and HIV-2 RNA copackaging

Coexpressed constructs ^a	CeFP ⁺ particles			RNA labeling efficiency ^c	
	Total no. analyzed	YFP ⁺ (%)	mCherry ⁺ (%)		
HIV-1–GagCeFP-MS2SL + HIV-2–DIS3GC-GagCeFP-BglSL					
Expt 1	10,420	35.0	45.9	12.4	93.3
Expt 2	7,558	42.6	38.6	12.2	93.4
Expt 3	5,866	47.9	32.5	11.3	91.7
Expt 4	8,644	47.6	33.1	12.5	93.2
Expt 5	9,474	49.7	33.1	9.6	92.4
Expt 6	11,034	39.2	37.5	13.6	90.3
Expt 7	11,664	33.4	45.6	12.6	91.6
HIV-1–DIS6C-GagCeFP-MS2SL + HIV-2–DIS6G-GagCeFP-BglSL					
Expt 1	2,762	26.2	34.2	31.9	92.3
Expt 2	2,853	24.1	40.7	28.6	93.4
Expt 3	2,896	28.7	33.4	31.9	94.0
Expt 4	6,605	36.6	26.4	31.0	94.0
Expt 5	6,647	36.0	25.0	33.0	94.0
Expt 6	6,387	32.4	29.2	32.3	93.9
Expt 7	5,095	37.7	26.6	28.5	92.8
Expt 8	4,806	35.2	29.3	28.5	93.0
Expt 9	5,702	36.3	26.2	31.1	93.6
HIV-1–DIS6C-GagCeFP-MS2SL + HIV-2–DIS6C-GagCeFP-BglSL					
Expt 1	3,085	35.4	38.7	16.9	91.0
Expt 2	3,374	36.7	37.3	17.7	91.7
Expt 3	3,259	35.0	40.7	15.1	90.8
Expt 4	6,100	35.7	40.6	14.4	90.7
Expt 5	4,209	39.8	35.2	16.3	91.3
Expt 6	4,609	41.0	33.7	15.3	90.0
Expt 7	8,688	34.1	42.2	14.8	91.1
Expt 8	9,626	33.2	42.5	16.6	92.3

^a Although not indicated, both Bgl-mCherry and MS2-YFP were coexpressed in all experiments.

^b Average \pm SD for HIV-1–GagCeFP-MS2SL + HIV-2–DIS3GC-GagCeFP-BglSL, 12.0 ± 1.3 ; average \pm SD for HIV-1–DIS6C-GagCeFP-MS2SL + HIV-2–DIS6G-GagCeFP-BglSL, 30.8 ± 1.8 ; average \pm SD for HIV-1–DIS6C-GagCeFP-MS2SL + HIV-2–DIS6C-GagCeFP-BglSL, 15.9 ± 1.2 .

^c Calculated by adding the values from the three columns under the CeFP⁺ heading.

that HIV-1 RNA was not packaged efficiently, and copackaging was not observed (Table 3).

To address the possibility that HIV-1 RNA may not have been expressed in the cells transfected with HIV-2 and therefore not packaged, we generated HIV-2–GagCeFP-BglSL-noTatRev, a construct that contains a frameshift mutation in the overlapping region of the HIV-2 *tat* and *rev* genes, to abolish the expression of functional Tat and Rev (Fig. 3B). Lacking functional Tat and Rev, HIV-2–GagCeFP-BglSL-noTatRev does not produce Gag and therefore does not generate viral particles (data not shown). However, when cotransfected with HIV-1–noGag-MS2SL that expressed functional Tat and Rev, HIV-2–GagCeFP-BglSL-noTatRev was able to produce viral particles. This strategy ensures the expression of HIV-1 RNA in the virus-producing cells. Analyses of these particles revealed abundant mCherry signals, indicating efficient packaging of HIV-2 RNA but very little YFP signals, indicating the lack of encapsidation of HIV-1 RNA (Table 3). To investigate whether HIV-1 RNA packaging can be rescued by base pairing in the DIS sequences with HIV-2 RNA, we generated HIV-1–DIS6C-noGag-MS2SL and HIV-2–DIS6G-GagCeFP-BglSL-noTatRev, two variants containing DIS sequences that, when both HIV-1 and HIV-2 Gag proteins were expressed, pro-

moted heterologous RNA packaging. Single-virion analyses revealed that the complementary DIS mutations were not sufficient to promote HIV-1 RNA packaging to detectable levels (Table 3).

Next, we examined the effects of eliminating HIV-2 Gag in heterologous RNA copackaging experiments. We generated HIV-2–noGag-MS2SL, which contains two inactivating mutations in the *gag* gene: a substitution mutation in the MA-encoding region that generated a stop codon and a frameshift in the CA-encoding region that disrupts the Gag reading frame. We then analyzed particles generated by coexpressing HIV-1–GagCeFP-MS2SL, HIV-2–noGag-MS2SL, MS2-YFP, and Bgl-mCherry and observed a heterologous RNA copackaging frequency of $\sim 10\%$ (range, 9% to 11%) (Table 4), which is similar to the frequency observed when both HIV-1 and HIV-2 Gag were present (Table 1). We then generated HIV-2 constructs that contained mutations in DIS sequences (Fig. 3C) and examined whether these mutations had the same effects on RNA copackaging when only HIV-1 Gag was present in the system. Along with MS2-YFP and Bgl-mCherry, coexpression of HIV-1–DIS6C–GagCeFP-MS2SL and HIV-2–DIS6C-noGag-BglSL resulted in $\sim 16\%$ heterologous RNA copackaging (Table 4), whereas coexpression of HIV-1–

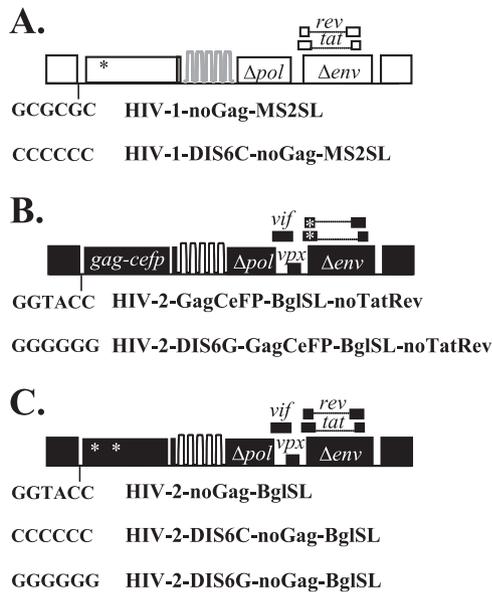


FIG. 3. General structures of constructs used to examine the roles of HIV-1 and HIV-2 Gag proteins in heterologous RNA copackaging. (A) HIV-1 constructs that do not express functional Gag. These constructs harbor a frameshift mutation in *gag*. (B) HIV-2 constructs that do not express functional Tat and Rev. These constructs harbor a frameshift mutation that simultaneously prevents expression of functional HIV-2 Tat and that of functional Rev. (C) HIV-2 constructs that do not express functional Gag and their DIS sequences. Each construct contains two mutations in *gag* to prevent expression of functional Gag. DIS sequences of the constructs are indicated. Frameshift mutations are indicated by asterisks.

DIS6C–GagCeFP-MS2SL and HIV-2–DIS6G-noGag-BglSL resulted in ~39% RNA copackaging (Table 4). The introduction of DIS mutations in the absence of HIV-2 Gag influenced copackaging in a manner parallel to that observed when both HIV-1 and HIV-2 Gag were present. Therefore, HIV-1 Gag alone is most likely responsible for HIV-1/HIV-2 RNA copackaging; although coassembly of HIV-1 and HIV-2 Gag occurs frequently, this event is not driving heterologous RNA copackaging.

DISCUSSION

During the assembly process of all known retroviruses, Gag proteins select from a large pool of total cellular RNAs and package two copies of full-length, single-stranded viral RNA into viral particles. In some viruses, such as HIV-1 and HIV-2, RNA selects its copackaged partner (or dimerizes) prior to encapsidation (4, 23, 27). Considering the specificity of this process, the RNA-RNA and RNA-protein interactions involved are likely to be highly regulated and coordinated events. The packaging of two distinct retroviral RNAs into the same virus particle requires at least a partial overlap, spatially as well as temporally, of the two RNAs in the producer cell, as well as similar packaging determinants and mechanisms of the two viruses. The work described in this report explored the factors that influence the copackaging of two divergent human retroviruses, HIV-1 and HIV-2. The copackaging of the heterologous RNAs is a determinant for potential genetic interactions;

TABLE 3. Single-virion analyses of RNAs packaged by HIV-2 Gag

Coexpressed constructs ^a	CeFP ⁺ particles			RNA labeling efficiency ^c
	Total no. analyzed	YFP ⁺ (%)	mCherry ⁺ (%)	
HIV-1–noGagCeFP-MS2SL + HIV-2–GagCeFP-BglSL				
Expt 1	2,405	0.0	92.6	0.9
Expt 2	835	0.0	96.3	0.2
Expt 3	1,227	0.0	92.3	2.1
Expt 4	1,089	0.0	93.8	1.0
Expt 5	909	0.0	97.4	0.9
Expt 6	614	0.0	98.4	0.0
Expt 7	1,548	0.0	91.5	1.2
HIV-1–noGagCeFP-MS2SL + HIV-2–GagCeFP-BglSL-noTatRev				
Expt 1	495	0.0	97.0	0.6
Expt 2	428	0.0	93.9	1.4
Expt 3	391	0.0	98.0	0.3
Expt 4	1,176	0.0	90.3	0.7
HIV-1–DIS6C-noGagCeFP-MS2SL + HIV-2–DIS6G-GagCeFP-BglSL-noTatRev				
Expt 1	687	0.0	91.7	0.9
Expt 2	527	0.0	91.7	0.8
Expt 3	1,146	0.0	94.9	1.6
Expt 4	810	0.0	90.6	1.5

^a Although not indicated, both Bgl-mCherry and MS2-YFP were coexpressed in all experiments.
^b Average ± SD for HIV-1–noGagCeFP-MS2SL + HIV-2–GagCeFP-BglSL, 0.9 ± 0.7; average ± SD for HIV-1–noGagCeFP-MS2SL + HIV-2–GagCeFP-BglSL-noTatRev, 0.8 ± 0.5; average ± SD for HIV-1–DIS6C-noGagCeFP-MS2SL + HIV-2–DIS6G-GagCeFP-BglSL-noTatRev, 1.2 ± 0.4.
^c Calculated by adding the values from the three columns under the CeFP⁺ heading.

TABLE 4. Single-virion analyses of RNAs packaged by HIV-1 Gag

Coexpressed constructs ^a	CeFP ⁺ particles			RNA labeling efficiency ^c	
	Total no. analyzed	YFP ⁺ (%)	mCherry ⁺ (%)		YFP ⁺ + mCherry ⁺ (%) ^b
HIV-1-GagCeFP-MS2SL + HIV-2-noGagCeFP-BglSL					
Expt 1	1,243	46.3	33.8	10.1	90.2
Expt 2	349	22.6	57.9	9.5	90.0
Expt 3	1,399	45.6	36.7	10.1	92.4
Expt 4	1,405	47.0	35.9	9.1	92.0
Expt 5	1,240	42.5	40.9	9.0	92.4
Expt 6	2,839	27.2	53.9	8.9	90.0
Expt 7	2,376	45.5	37.7	9.9	93.1
Expt 8	4,052	27.8	54.8	9.7	92.3
Expt 9	2,016	47.2	37.1	9.0	93.3
Expt 10	4,701	53.9	26.9	10.3	91.1
Expt 11	5,218	58.0	23.7	10.5	92.2
Expt 12	6,564	48.9	31.7	10.5	91.1
Expt 13	6,789	42.7	38.7	9.3	90.7
HIV-1-DIS6C-GagCeFP-MS2SL + HIV-2-DIS6C-noGagCeFP-BglSL					
Expt 1	3,152	45.4	34.1	12.7	92.2
Expt 2	4,006	47.8	30.3	15.4	93.5
Expt 3	5,282	48.0	29.2	17.5	94.7
Expt 4	5,703	45.6	33.1	15.8	94.5
Expt 5	6,045	44.7	27.5	18.5	90.7
HIV-1-DIS6C-GagCeFP-MS2SLL + HIV-2-DIS6G-noGagCeFP-BglSL					
Expt 1	3,881	29.4	28.7	36.7	94.8
Expt 2	4,589	33.7	21.5	39.8	95.0
Expt 3	5,334	34.4	22.7	38.3	95.4
Expt 4	5,020	29.6	26.1	38.8	94.5
Expt 5	4,862	24.8	28.5	41.1	94.4

^a Although not indicated, both Bgl-mCherry and MS2-YFP were coexpressed in all experiments.

^b Average \pm SD for HIV-1-GagCeFP-MS2SL + HIV-2-noGagCeFP-BglSL, 9.7 ± 0.6 ; average \pm SD for HIV-1-DIS6C-GagCeFP-MS2SL + HIV-2-DIS6C-noGagCeFP-BglSL, 16.0 ± 2.2 ; average \pm SD for HIV-1-DIS6C-GagCeFP-MS2SLL + HIV-2-DIS6G-noGagCeFP-BglSL, 38.9 ± 1.6 .

^c Calculated by adding the values from the three columns under the CeFP⁺ heading.

furthermore, this system allowed us to study the mechanisms of retroviral RNA dimerization and packaging.

In this study we found that, within a population of particles containing similar proportions of HIV-1 and HIV-2 RNAs, about 10% of the particles contained both HIV-1 and HIV-2 RNAs. Equal expression and random assortment of RNAs from two proviruses predict that 50% of the virions would be heterozygous particles, containing one RNA derived from each provirus. The 10% copackaging frequency implies that HIV-1 and HIV-2 RNA packaging mechanisms have sufficient overlap to yield a modest but reproducible level of copackaging but are divergent enough that HIV-1 and HIV-2 RNAs still prefer to homodimerize. We have previously measured recombination between HIV-1 and HIV-2 using two modified genomes, each carrying a mutated green fluorescent protein (*gfp*) gene; recombination between the two mutated genes could reconstitute a functional *gfp* gene, and its expression could be measured. We found that a very small percentage ($\sim 0.5\%$) of infected cells had a GFP⁺ phenotype (25), which is much lower than that generated between two HIV-1 or two HIV-2 viruses ($\sim 7\%$) (5, 29). In all three studies, the *gfp* gene was used as a target sequence for reporting recombination; thus, regardless of the sequence diversity elsewhere in the viral genomes, the target regions between the mutations in the *gfp* genes contain

identical sequences. Results from this study indicate that $\sim 10\%$ of the particles have copackaged HIV-1 and HIV-2 RNAs and that the low copackaging frequency was a contributing factor but not the only one that caused the lower recombination rate. Together, these studies revealed that there is a block or barrier in the reverse transcription step that limits the completion of DNA synthesis or recombination in the target *gfp* gene between these heterologous genomes. The details of the mechanism for this block or barrier are currently unknown. It is possible that the two copackaged RNAs have different structures, thereby inhibiting DNA synthesis or preventing RT from switching templates; alternatively, it is also possible that the low sequence homology flanking the *gfp* genes reduces RT template switching within the *gfp* gene. Further experimentation is needed to distinguish between these possibilities.

In this report, we studied factors that may restrict the heterodimerization of HIV-1 and HIV-2 RNA and thus restrict copackaging. We found that changing the DIS to the same palindrome (GCGCGC) to allow the DIS sequences of HIV-1 and HIV-2 RNA to form base pairs did not increase the copackaging, most likely because the two RNAs still homodimerize efficiently. The largest improvement of heterologous RNA copackaging was observed when we simultaneously encouraged RNA heterodimerization and discouraged homodimerization by changing

the HIV-1 DIS to CCCCCC and the HIV-2 DIS to GGGGGG. These results not only revealed that DIS sequences can mediate dimerization of the heterologous RNAs but also shed light on the mechanisms of Gag-RNA interactions.

Although RNA partner selection occurs prior to the encapsidation of HIV-1 or HIV-2 RNA, it is unclear how HIV Gag proteins distinguish between monomeric and dimeric RNAs. Studies of MLV packaging showed that the monomeric and dimeric RNAs have very different secondary structures (10, 15, 21). Furthermore, dimerization of MLV RNAs exposes high-affinity binding sites for nucleocapsids, which is critical for RNA packaging (16, 22). It is thought that exposure of these sites allows Gag binding, leading to the incorporation of RNA into virus particles. Despite its attractiveness, this model has been difficult to extend to the RNA packaging mechanisms of HIV-1, mainly because structural analyses of HIV-1 RNA have not revealed a drastic difference between monomeric RNA and dimeric RNA (35); hence, the relationship between switching the RNA states and RNA-Gag molecular recognition is not well defined. Our study revealed surprising insights into this process. Despite the low homology of HIV-1 and HIV-2 leader sequences, when DIS sequences were manipulated to form base pairing between these two heterologous RNAs but not between homologous RNAs, we observed increased copackaging. If the HIV-1 and MLV Gag-RNA recognition mechanisms are similar, then our results suggest that base pairing of the DIS between these two heterologous RNAs can expose the hidden high-affinity binding sites recognized by HIV-1 Gag. The limited homology between the 5' leader sequences of HIV-1 and HIV-2 makes it less likely that there are many other conserved interactions between these two RNAs; hence, this hypothesis implies that base pairing of DIS provides a molecular switch that directly or indirectly exposes Gag-binding sites.

A possible model for the mechanism of heterologous RNA copackaging is that such events are driven by the coassembly of Gag polyproteins from different viruses. However, our results indicated that HIV-1 Gag, in the absence of HIV-2 Gag, can mediate heterologous RNA copackaging at levels similar to those observed with both HIV-1 and HIV-2 Gag proteins. This observation defines the *trans*-acting element required for the heterologous RNA copackaging, that is, the Gag protein that recognizes its own RNA and the heterologous RNA.

Replication strategies used by retroviruses ensure frequent generation of recombinants, as one viral particle has two full-length viral genomes and RT switches templates during DNA synthesis. Although copackaged RNAs are derived from the same types of viruses most of the time, there are infrequent occurrences of heterologous RNA copackaging, which provides the backdrop for the generation of chimeric viruses from two distantly related viruses. Delineating the requirements and factors that influence heterologous RNA copackaging allows us to understand the potential for the genetic interactions between distantly related viruses and also provide insights into the general mechanisms of RNA-RNA and RNA-Gag interactions that lead to encapsidation of viral genomes.

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