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Mannose-rich glycosylation patterns on HIV-1 subtype C gp120 and sensitivity to the lectins, Griffithsin, Cyanovirin-N and Scytovirin

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Abstract

Griffithsin (GRFT), Cyanovirin-N (CV-N) and Scytovirin (SVN) are lectins that inhibit HIV-1 infection by binding to multiple mannose-rich glycans on the HIV-1 envelope glycoproteins (Env). Here we show that these lectins neutralize subtype C primary virus isolates in addition to Env-pseudotyped viruses obtained from plasma and cervical vaginal lavages. Among 15 subtype C pseudoviruses, the median IC₅₀ values were 0.4, 1.8 and 20.1 nM for GRFT, CV-N and SVN, respectively, similar to what was found for subtype B and A. Analysis of Env sequences suggested that concomitant lack of glycans at positions 234 and 295 resulted in natural resistance to these compounds, which was confirmed by site-directed mutagenesis. Furthermore, the binding sites for these lectins overlapped that of the 2G12 monoclonal antibody epitope, which is generally absent on subtype C Env. This data support further research on these lectins as potential microbicides in the context of HIV-1 subtype C infection.

Keywords

Griffithsin; Cyanovirin-N; Scytovirin; lectins; HIV-1 subtype C; microbicides; 2G12 monoclonal antibody

INTRODUCTION

Griffithsin (GRFT) is a lectin isolated from the red algae *Griffithsia* sp. found in the coastal waters off New Zealand (Mori et al., 2005). It is a 121 amino acid dimeric protein with a domain swapped structure (Ziolkowska et al., 2006). This lectin neutralizes HIV-1 by binding to mannose-rich glycans found on the envelope glycoproteins. Both native and recombinant GRFT display potent antiviral activities against laboratory adapted strains and primary isolates of M- and T- tropic HIV-1. This compound was also shown to be active against a broad range of HIV-1 including 4 subtype C viruses (O'Keefe et al., 2009). A second lectin, Cyanovirin-N (CV-N) isolated much earlier from the blue green algae *Nostoc*

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elliposporum (Boyd et al., 1997) is a 101 amino acid protein that has been shown to exist either as a quasi-symmetric two-domain monomer or as a domain-swapped dimer (Barrientos et al., 2002; Botos and Wlodawer, 2005). Like GRFT, CV-N binds to mannose-rich glycans and both native and recombinant CV-N have shown potent anti-HIV-1 activity *in vitro* (Bolmstedt et al., 2001; Esser et al., 1999). A third protein, Scytovirin (SVN) is a 95 amino acid lectin isolated from the cyanobacterium *Scytonema varium* (Bokesch et al., 2003). SVN is expressed as a single amino acid chain with extensive internal sequence duplication. This protein is found exclusively as a monomer and has been shown to neutralize both laboratory adapted strains and primary isolates of HIV-1 by interacting with mannose-rich glycans on the viral envelope (Moulaei et al., 2007; Xiong et al., 2006; Ziolkowska and Wlodawer, 2006).

As a result of their ability to block HIV-1 entry *in vitro*, GRFT, CV-N and SVN have been proposed as potential microbicides to prevent the sexual transmission of HIV-1. Although these compounds are not HIV-specific, they target high-mannose arrays that are present on all HIV envelope glycoproteins. Since such arrays are uncommon in mammalian cells, these compounds are not likely to be toxic to human cells *in vivo* even at relatively high concentrations (Balzarini, 2005; Balzarini and Van Damme, 2007). Furthermore, a recombinant GRFT produced in the tobacco-like plant *Nicotiana benthamiana* was shown to be non-toxic in a rabbit vaginal irritancy model and in human cervical explants (O'Keefe et al., 2009). Although GRFT, CV-N and SVN have not yet been tested in human clinical trials it is noteworthy that CV-N was shown to be effective in protecting pigtailed macaques after vaginal and rectal challenges with high dose of SHIV 86.9P (Tsai et al., 2004; Tsai et al., 2003).

HIV-1 can develop resistance to CV-N by partial deglycosylation, specifically deleting glycans at positions 230, 289, 295, 332, 339, 386, 392 and 448 in HIV-IIIB (Balzarini et al., 2006; Hu, Mahmood, and Shattock, 2007). This suggests that resistance to CV-N may be hard to generate as it will require multiple mutations in the viral genome although additional studies using primary viruses are needed. Some of these glycans also form part of the epitope for the broadly neutralizing monoclonal antibody (mAb) 2G12. This antibody directly binds glycans at positions 332, 339 and 392 (Calarese et al., 2003), while glycans at positions 295, 386 and 448 though not directly involved in this epitope, may influence its conformation (Sanders et al., 2002; Scanlan et al., 2002). Subtype C viruses are commonly resistant to 2G12 neutralization. This has been attributed to their frequent lack of the 295 glycosylation site although re-introduction of this glycan only partially restores sensitivity to this mAb (Binley et al., 2004; Chen et al., 2005; Gray et al., 2007b). Although it is known that GRFT and SVN bind to mannose-rich glycans found on HIV-1 envelope, the specific glycans involved in this binding have not yet been identified.

Despite subtype C viruses being responsible for over 50% of global infections (<http://www.unaids.org>) many of the antiviral studies with GRFT, CV-N and SVN have been done on subtype B viruses. Significant differences exist in the pattern of glycosylation between subtype B and C envelope glycoproteins (Zhang et al., 2004). For example, about 80% of subtype B envelopes are glycosylated at position 295 while only ~20% of subtype C viruses are glycosylated at the same position. The opposite pattern is observed at position 230 where over 70% of subtype C envelopes are glycosylated compared to only 20% of subtype B. It is known that differences between subtypes can affect their susceptibility to neutralizing agents as for the 2G12 mAb described above (Binley et al., 2004; Gray et al., 2006). Therefore, we aimed to examine the sensitivity of a large collection of subtype C viruses to GRFT, CV-N and SVN and compare these to subtypes B and A strains. Furthermore, we examined the role of the 2G12 epitope in these lectin's binding sites.

RESULTS

GRFT, CV-N and SVN neutralize HIV-1 subtype C isolates from blood and cervico-vaginal lavages

To determine the ability of GRFT, CV-N and SVN to inhibit HIV-1 subtype C infection of peripheral blood mononuclear cells (PBMC), we performed neutralization assays using blood-derived subtype C primary isolates Du151, COT9 and Du179. Viral infection was measured by p24 ELISA after 4 days of culture. The concentration of lectin that inhibited 80% of virus infection (IC_{80}) is reported. All three lectins neutralized these viruses in adose-dependent manner (Figure 1). The IC_{80} values were similar for all viruses with GRFT and CV-N being more potent than SVN. The mean IC_{80} values for all 3 viruses were 42.7 ± 4.4 , 77.0 ± 18.2 and 368.4 ± 22.7 nM for GRFT, CV-N and SVN, respectively.

Next we tested the ability of GRFT, CV-N and SVN to inhibit 50% (IC_{50}) HIV-1 infection in the TZM-bl cell-line based neutralization assay (Montefiori, 2004). For this we used 11 subtype C Env-pseudotyped viruses cloned from plasma of adult patients with acute or early HIV infection and from pediatric patients during chronic infection (see Methods for details). For comparison we included 6 viruses from the subtype B panel described by Li *et al.* (Li *et al.*, 2005) and 5 subtype A viruses from acutely infected individuals (Long *et al.*, 2002; Neilson *et al.*, 1999). All viruses used the CCR5 coreceptor. GRFT and CV-N potently neutralized most subtype C viruses with a median IC_{50} of 0.4 and 1.2 nM, respectively, (Figure 2) which were not significantly different. SVN was the least effective with a median IC_{50} of 20.1 nM. The single-cycle pseudovirus assay required considerably lower concentrations of lectin to inhibit viral infection compared to the PBMC assay even when comparing IC_{80} values (not shown). Nevertheless, the trends were the same in both assays as was noted for Du151.2 and COT9.6 pseudoviruses compared to their matched primary viruses in PBMC (Table 1 and Figure 1). There were no significant differences in mean IC_{50} values between subtypes C, B and A viruses for all 3 lectins ($p > 0.05$) (Table 1). However, 2 subtype C viruses (CAP206.08J and CAP63.A9J) and one subtype B virus (CAAN5342.A2) showed unusually high IC_{50} values suggesting natural resistance to these compounds (Figure 2). We also found a significant positive correlation ($p < 0.05$) between GRFT and CV-N sensitivity for subtype C, and between CV-N and SVN for both subtype C and B Env-pseudotyped viruses (data not shown).

We also tested the sensitivity of four subtype C Env isolated from cervical-vaginal lavages (CVL) of HIV-1 positive women during acute infection (Bronwen Lambson, unpublished data). These viruses also showed varying sensitivities to the lectins with one (CAP177.cv196) showing a high level of resistance to both GRFT and CV-N (Figure 3). For CAP63.cv16GA a matched blood sample was also available and the CVL and plasma derived viruses showed similar sensitivities to the three lectins (Table 1). In general, viruses from CVL were inhibited by GRFT, CV-N and SVN with IC_{50} in the nanomolar range similar to those derived from plasma.

Correlation between HIV-1 mannose-rich glycosylation patterns and sensitivity to GRFT, CV-N and SVN

Given that these lectins bind mannose-rich glycans, we analyzed the predicted glycosylation patterns of all the viruses used in our neutralization assays to determine if there was any correlation with their sensitivity to GRFT, CV-N and SVN. This analysis included all 11 predicted mannose-rich sites on HIV-1 gp120 some of which are involved in mAb 2G12 binding (labeled with asterisks in Table 1). Significant differences were noted between subtypes as previously reported (Zhang *et al.*, 2004). In this study, 13 of the 15 subtype C viruses lacked the 295 glycan compared to more than half of the subtype B and A viruses

where this glycan was intact. On average, viruses from subtypes C and B lacked 2 of the 11 predicted glycans although there was a range with some viruses lacking as many as 5 sites. The subtype A viruses had a larger number of missing sites and they all lacked the 230 and 289 glycans. Only one virus was predicted to have all 11 glycans, a subtype C virus (CAP270.cv144) isolated from CVL.

The five viruses (four subtype C, including two from CVL, and one subtype B) that were the least sensitive to GRFT lacked both the 234 and 295 glycosylation sites. Two of these five viruses (CAP206.08J and CAAN5342.A2) were also the least sensitive to CV-N and SVN and a third (CAP177.cv196) also required above average levels of CV-N for inhibition. We concluded that the absence of both the 234 and 295 glycosylation sites was responsible for this decreased sensitivity to these lectins. However, the lack of either of these two glycans in isolation did not affect sensitivity. Thus WITO4160.33 and Q842.d12 which lacked the 234 glycan but had the 295 glycan showed similar sensitivity to viruses with both glycans. Similarly viruses that lacked the 295 glycan but retained the glycan at position 234 (most subtype C viruses) were not unusually resistant. Overall there was no correlation between the number of missing glycans and resistance to these lectins. The fewer mannose-rich glycosylation sites on subtype A Envs did not reduce their sensitivity.

The 234 and 295 glycosylation sites are involved in GRFT, CV-N and SVN neutralization of HIV-1

In order to examine the contribution of the glycan at position 295 to lectin neutralization, we introduced this glycosylation site in 3 viruses, Du151.2, COT9.6 and COT6.15, that already had the 234 glycan (see Table 1). For all three viruses the introduction of the 295 glycosylation site further increased their sensitivity to GRFT by 7 to 50 fold (Table 2). However, the addition of the 295 glycan had no effect on their sensitivity to CV-N and SVN suggesting that this glycan is essential for GRFT binding only. As GRFT is more promiscuous in its binding to high-mannose oligosaccharides than either SVN or CV-N (which only bind to oligomannose-8 or -9) this could also indicate that the glycan at position 295 is a smaller oligosaccharide such as oligomannose-5 or -7.

We also introduced both the 234 and 295 glycosylation sites into 3 viruses, CAP206.08J, CAP63.A9J and CAAN5342.A2 that lacked both of these sites and were the most resistant to the lectins, to determine if this would increase their sensitivity. Reconstruction of the 234 glycosylation site in both CAP206.08J and CAP63.A9J reduced virus infectivity of TZMbl-cells by ~2-fold. We were unable to determine whether this was due to a decrease in the production of viral particles in 293T cells or a defect in the virus's capacity to infect cells. However, the introduction of 295 and to a lesser extent 234, increased the sensitivity of all three viruses to GRFT (Figure 4A). Furthermore, when these two glycans were introduced simultaneously there was an additional increase in sensitivity. The introduction of the 234 and 295 glycosylation sites also had a similar effect on CV-N sensitivity, with the exception of CAP63.A9J where addition of the 234 glycosylation site had no effect (Figure 4B). For SVN, the 234 glycosylation site had a more significant impact on sensitivity than the 295 although for CAP63.A9J addition of a glycan at position 295 actually increased sensitivity above the wild-type (Figure 4C). Collectively, these data suggested that the 234 and 295 glycosylation sites are involved in lectin binding to the HIV-1 envelope glycoproteins and subsequent virus neutralization, although for CV-N and SVN their involvement may be strain-dependent.

GRFT, CV-N and SVN binding sites overlap the 2G12 epitope

Since five of the 11 mannose-rich sites on gp120 form the 2G12 mAb epitope, we investigated whether lectin binding interfered with access to this epitope. We used a

competition ELISA to measure the amount of p24 antigen captured onto 2G12-coated plates after pre-incubation of the virus with GRFT, CV-N or SVN. The 2G12 mAb was competed against itself as a positive control. We tested the subtype B virus QH0692.42 which is sensitive to 2G12 with an IC_{50} of 2.8 $\mu\text{g/ml}$ (Li et al., 2005). Subtype C viruses are generally insensitive to 2G12 and so we used modified subtype C gp160 where the 2G12 epitope had been partially reconstituted (Gray et al., 2007b). Thus, for COT6.15 the addition of glycans at positions 295 and 448 (COT6-V295N/S448N) increased sensitivity from $IC_{50} >100 \mu\text{g/ml}$ to 64.5 $\mu\text{g/ml}$. There was also increased binding of the 2G12 mAb to this mutant (Gray et al., 2007b). For COT9.6 the addition of glycans at positions 295 and 442 (COT9-V295N/K442N) increased the sensitivity to 2G12 from an IC_{50} of 68.7 $\mu\text{g/ml}$ to 33.9 $\mu\text{g/ml}$.

Pre-incubation of the 3 viruses with each lectin significantly reduced the amount of virus captured onto 2G12-coated plates suggesting competition between these compounds (Figure 5). Similar to what was seen in the neutralization assays; GRFT and CV-N were more potent than SVN at inhibiting virus capture. The effect was dose-dependent although higher amounts of lectin were required to inhibit virus capture compared to neutralization. However, for all 3 viruses, virus capture by 2G12 was inhibited at equivalent lectin concentrations despite the 10–20 fold difference in sensitivities to 2G12 neutralization. Together with the mutagenesis results (Table 2) these data suggested an overlap between the 2G12 epitope and GRFT, CV-N and SVN binding sites.

DISCUSSION

In this study we have shown that HIV-1 subtype C viruses, including those from the vaginal tract are sensitive to the mannose-binding lectins, GRFT, CV-N and SVN. Despite subtype-specific differences in mannosylation patterns, their sensitivity was similar to what has been previously reported for subtype B viruses (Bokesch et al., 2003; Boyd et al., 1997; Mori et al., 2005). GRFT and CV-N were more potent than SVN which required higher concentrations to prevent HIV-1 entry. We identified glycans at positions 234 and 295, in addition to the 2G12 mAb epitope, as playing an important role in the binding of these lectins. Our data support the continued development of GRFT, CV-N and SVN as potential microbicides since these compounds are likely to be effective against HIV-1 subtype C mucosal transmissions.

Inhibition of HIV-1 subtype C infection by GRFT, CV-N and SVN in both primary cells and a cell line were similar although higher concentrations were required for PBMC neutralization. The increased sensitivity of TZM-bl cells compared to PBMC is consistent with studies of antibody neutralization, where single-cycle pseudovirus assays are generally more sensitive than PBMC infectivity assays (Fenyo et al., 2009). The longer duration of culture and multiple replication cycles in the PBMC assay probably account for this. Unlike Huskens and colleagues, we did not find any enhanced infectivity of HIV-1 in PBMC treated with CV-N, indicating that CV-N was not mitogenic in our hands (Huskens et al., 2008). The greater neutralization potency of GRFT and CV-N compared to SVN may be due to the fact that a GRFT dimer has six mannose binding sites and a CV-N dimer has four binding sites. In contrast, the SVN monomer has only two mannose residue binding sites (Moulaei et al., 2007; Ziolkowska et al., 2006; Ziolkowska and Wlodawer, 2006). The correlation between IC_{50} values of the compounds within certain subtypes suggested that these lectins share some binding sites, a finding which we confirmed for a few loci. Although, we have shown efficacy of these compounds against subtype C viruses in primary PBMC, it will be important to test the anti-HIV activity of GRFT, CV-N and SVN using dendritic cells and macrophages as these cells play an important role during the sexual

transmission of HIV (Lederman, Offord, and Hartley, 2006; Pohlmann, Baribaud, and Doms, 2001).

Subtype C viruses isolated from both adult and pediatric HIV infections as well as those from acute and chronic infection were sensitive to these lectins. Our data are in line with previously reported inhibitory concentrations for subtype B viruses and confirm that these compounds are active against multiple subtypes (Bokesch et al., 2003; Boyd et al., 1997; Mori et al., 2005; O'Keefe et al., 2009). Given that CV-N and GRFT are being proposed as microbicides, we also considered it important to test them against viruses found in the vaginal tract. It has been reported that virus compartmentalization can occur during HIV infection raising the possibility of differential sensitivity of viruses from CVL compared to blood (Kemal et al., 2003). However, our data showed that CAP63.A9J from plasma and CAP63.cv16GA from CVL of the same HIV-positive woman had similar sensitivity to all three lectins. We attributed this to the fact that these two viruses have minimal genetic differences and the same mannose-rich glycosylation patterns (Table 1). Also these viruses are from early infection and, therefore, it may be too early for compartmentalization to develop. Nevertheless, further studies with more pairs of plasma and CVL viruses, from the same individual with a demonstrated virus compartmentalization, are needed to determine if viruses from CVL are as sensitive as those present in blood.

Despite differences in mannose-rich glycosylation patterns, subtype C, B and A viruses (Zhang et al., 2004) showed similar sensitivities to GRFT, CV-N, and SVN. This result was unexpected given that these compounds bind high-mannose residues. Analysis of the glycosylation patterns suggested that absence of glycans at positions 234 and 295 was associated with natural resistance to GRFT and in some cases to CV-N and SVN. Since subtype C viruses generally lack 295N it was surprising that these viruses were as sensitive to the lectins as subtype B and A viruses that generally retained this site. However, a concomitant lack of the 234 glycosylation site was also required to reduce sensitivity and since this glycan is present on ~80% of subtype C viruses, this may account for the comparable sensitivity with subtype B. The important role of these glycans in conferring sensitivity to these lectins was confirmed by site-directed mutagenesis. Interestingly, when the 295 or 234 glycosylation sites were individually reconstituted they had a significant impact in some viruses. This suggests that in natural Env there may be considerable redundancy or overlap in the glycans that are bound by these compounds. We did not find a correlation between the number of predicted mannose-rich glycans and sensitivity to the lectins, unlike an earlier study on CV-N in subtype B viruses (Balzarini et al., 2006). For example, CAP270.cv144 which had all predicted 11 mannose-rich glycans was less sensitive to GRFT than viruses that had fewer predicted glycans. In fact our data suggested that the location of the glycan was more important than the number of glycans. Thus, in addition to mannose-rich glycans there may be other factors in the viral envelope that determine the sensitivity to GRFT, CV-N and SVN. These factors may include the Env structure, that can result in glycans that occupy the same position being exposed differently, and the fact that not all potential sites are glycosylated (Go et al., 2009).

Three studies examining CV-N escape pathways using laboratory-adapted HIV-1 subtype B viruses selected *in vitro* showed that deletion of glycans caused resistance to CV-N (Balzarini et al., 2006; Hu, Mahmood, and Shattock, 2007; Witvrouw et al., 2005). The large number of glycans involved suggested that these compounds have a high genetic barrier to resistance. In this study, we showed that for subtype C primary viruses that lack the 295 glycan, the absence of a single glycan at position 234 could in some cases confer natural resistance to these lectins. However, it is important to bear in mind that generating resistant variants *in vitro* is likely to be different to identifying loci associated with natural resistance as we have done here. Furthermore, naturally-occurring resistant viruses may have

compensatory mutations that are isolate-specific or not commonly present. Indeed, glycan 234 may play an important functional role given its relative conservation in subtype C, and this is first study to report its involvement in CV-N resistance in both subtypes C and B. Further studies are needed to clearly elucidate the position and number of glycans involved in CV-N resistance in primary viruses. It will also be important to investigate the ability of HIV-1 to develop resistance to GRFT and SVN, which may help to identify other GRFT and SVN binding sites on the virus.

The ability of GRFT, CV-N and SVN to compete against the 2G12 MAb for binding to HIV-1 implicates the 2G12 epitope in the binding of these compounds to HIV-1. These data are in agreement with previous reports that glycans in the 2G12 epitope are involved in CV-N and GRFT binding to HIV-1 gp120 (Esser et al., 1999; Mori et al., 2005). However, our study is the first to demonstrate the involvement of the 2G12 epitope in SVN binding to HIV-1. Nevertheless, the involvement of this epitope in the activity of these lectins to subtype C viruses is unclear as these viruses are generally insensitive to the 2G12 mAb (Chen et al., 2005; Gray et al., 2007b; Sanders et al., 2002). The sensitivity of subtype C viruses to GRFT, CV-N and SVN is likely due to the fact that these lectins bind many more sites than those involved in the 2G12 epitope.

GRFT, CV-N and SVN are being actively pursued as microbicides (Bokesch et al., 2003; O'Keefe et al., 2009; Shattock and Moore, 2003; Xiong et al., 2006). The search for HIV microbicides is driven by the challenges encountered in developing an effective HIV vaccine (Johnston and Fauci, 2007) and the fact that the majority of HIV infections around the world are sexually transmitted (Stein, 2003). Potential microbicides include non-specific inhibitors such as buffering agents that inactivate viruses by maintaining the acidic pH in the vaginal tract and moderately specific inhibitors such as anionic polymers that neutralize the virus by binding to its positively charged envelope glycoproteins (Cutler and Justman, 2008). Anti-retroviral agents that target the reverse transcriptase enzyme are being actively researched and herald a new phase in prevention strategies. GRFT, CV-N and SVN, however, hold an advantage over the above-mentioned compounds by selectively targeting the virus to inhibit its entry into susceptible cells. These compounds are small proteins that can be readily and cheaply produced. Furthermore, commensal lactobacilli can be engineered to produce these compounds which could be used to colonize vaginal mucosa and create an environment hostile to HIV-1 (Pusch et al., 2005). Since these compounds are not being developed as therapeutics, their exclusive use as prevention agents would avert the complications that may arise with the use of ARV as microbicides. In conclusion, our data support the continued development of GRFT, CV-N and SVN as microbicides against HIV-1 subtype C mucosal transmissions.

Materials and Methods

Viruses, gp160 envelope (Env) clones, MAbs, cell lines and lectins

HIV-1 subtype C isolates Du151, and Du179 were isolated in South Africa from individuals infected with HIV-1 subtype C (van Harmelen et al., 2001). COT9 is a South African pediatric isolate (Choge et al., 2006). HIV-1 subtype C Env clones Du151.2, CAP45.G3J, CAP210.E8J, CAP206.08J, CAP63.A9J, CAP228.51J and CAP256.F7J were amplified from South African individuals at the acute or early stage of HIV infection (Gray et al., 2007a; Li et al., 2006). HIV-1 subtype B Env, CAAN5342.A2, WITO4160.33, THRO4156.18, AC10.0.29, QH0692.42 and PVO.4 were amplified from acutely infected individuals from the U.S.A, Trinidad and Tobago and Italy (Li et al., 2005). Subtype C Env COT9.6, COT6.15, RP6.6 and TM7.9 were derived from South African pediatric HIV-1 isolates. CAP63.cv16GA, CAP177.cv196, CAP261.cv193 and CAP270.cv144 were amplified from cervico-vaginal lavages (CVL) obtained during the acute phase of infection from HIV

positive women in the CAPRISA cohort in South Africa. The pSG3Δ*env* plasmid was obtained from Beatrice Hahn. The mAb 2G12 was obtained from the NIH Reference and Reagent Program and the IAVI Neutralizing Antibody Consortium. The TZM-bl cell line was obtained from the NIH Reference and Reagent Program (Cat No 8129) and the 293T cell line was obtained from the American Type Culture Collection. Both cell lines were cultured in DMEM containing 10% fetal bovine serum (FBS). Cell monolayers were disrupted at confluence by treatment with 0.25% trypsin in 1 mM EDTA. Recombinant GRFT, CV-N, and SVN purified from *E.coli* were provided by Dr. Barry O'Keefe from NCI.

HIV-1 neutralization assay in peripheral blood mononuclear cells

The neutralization assay in PBMC was carried out as described by Bures *et al.* (Bures et al., 2000). Briefly, a three-fold dilution series of GRFT, CV-N, and SVN in 40 μL of RPMI 1640 containing 20% FBS and interleukin-2 (growth medium) was prepared in triplicate in a U-bottom 96-well plate. Five hundred TCID₅₀ of HIV-1 primary isolate in 15 μL of growth medium was added to each well and the plate was incubated at 37°C for 1 hour. Then 100 μL of 5 × 10⁶ cells/mL phytohemagglutinin/interleukin-2 stimulated PBMC (PHA-PBMCs) was added to each well. The following day cells were washed 3 times with RPMI 1640 with 20% FBS and resuspended in 155 μL of fresh growth medium. The culture supernatant was collected twice daily and replaced with an equal amount of fresh growth medium. For each harvest the p24 antigen concentration in the virus control wells was measured by ELISA using the Vironostika HIV-1 Antigen Microelisa System (Biomerieux, Boesind, the Netherlands), according to the manufacturer's instructions. The inhibitory activity of the lectins were measured at the time-point that corresponded to the early part of the linear growth period of the virus control (Zhou and Montefiori, 1997). The IC₈₀ were calculated by plotting the lectin concentration vs. the percentage inhibition in a linear regression using GraphPad Prism 4.0.

Generation of Env-pseudotyped virus stock

HIV-1 pseudoviruses were generated by co-transfection of the Env and pSG3Δ*env* plasmids (Wei et al., 2003) into 293T cells using the Fugene transfection reagent (Roche Applied Science, Indianapolis, IN). The TCID₅₀ of each virus stock was quantified by infecting TZM-bl cells with serial 5-fold dilutions of the supernatant in quadruplicate in the presence of DEAE dextran (37.5 μg/mL) (Sigma-Aldrich, St. Louis, MO). The Bright Glo™ Reagent (Promega, Madison, WI) was used to measure infection after 48 hours of tissue culture, according to the manufacturer's instructions. Luminescence was measured in a Wallac 1420 Victor Multilabel Counter (Perkin-Elmer, Norwalk, CT). The TCID₅₀ was calculated as described elsewhere (Johnson and Byington, 1990).

Single cycle neutralization assay (TZM-bl assay)

The pseudovirus neutralization assay was performed as described elsewhere (Montefiori, 2004). Briefly, three-fold dilution series of each lectin in 100 μL of DMEM with 10% FBS (growth medium) were prepared in a 96-well plate in duplicate. Two hundred TCID₅₀ of pseudovirus in 50 μL of growth medium was added and the mixture was incubated for 1 hour at 37°C. Then 100 μL of TZM-bl cells at a concentration of 1 × 10⁵ cells/mL containing 37.5 μg/mL of DEAE dextran was added to each well and cultured at 37°C for 48 hours. Infection was evaluated by measuring the activity of the firefly luciferase. Titers were calculated as the inhibitory concentration that causes 50% reduction (IC₅₀) of relative light unit (RLU) compared to the virus control (wells with no inhibitor) after the subtraction of the background (wells without both the virus and the inhibitor).

HIV-1 virion capture assay

High binding 96-well plates (Corning Incorporated Corning, New York, U.S.A) were coated overnight with 1 µg/well of the mAb 2G12 in NaHCO₃ (pH 8.5). The coated plates were washed three times with phosphate buffered saline (PBS) and blocked with 3% bovine serum albumin (BSA) in PBS at 37°C for 2 hours. Pseudoviruses were pre-incubated for an hour with different concentrations of 2G12, GRFT, CV-N, or SVN. After discarding the blocking solution the virus was added to the plate and left at 37°C for 2 hours. The plate was then washed three times with PBS and captured viruses were lysed with 150 µL of 0.5% Triton-X 100. The amount of p24 was measured using the Vironostika HIV-1 Antigen Microelisa System (Biomerieux, Boesend, the Netherlands), according to the manufacturer's instructions.

Site-directed mutagenesis

Putative glycosylation sites were introduced in HIV-1 gp120 using the QuikChange Site Directed Mutagenesis Kit (Stratagene, LaJolla, CA). The presence of the mutation was confirmed by sequencing using the ABI PRISM Big Dye Terminator Cycle Sequencing Ready Reaction kit (Applied Biosystem, Foster City, CA) and resolved on the ABI 3100 automated genetic analyzer.

Statistical analysis

The Mann-Whitney t-test was used to compare the median IC₅₀ of each lectin for the neutralization of subtype C, B and A viruses and the Wilcoxon matched pairs test was used to compare the median IC₅₀ of GRFT and CV-N for the neutralization of subtype C. GraphPad Prism 4.0 was used for the Spearman's nonparametric rank test. p-values of 0.05 were considered statistically significant.

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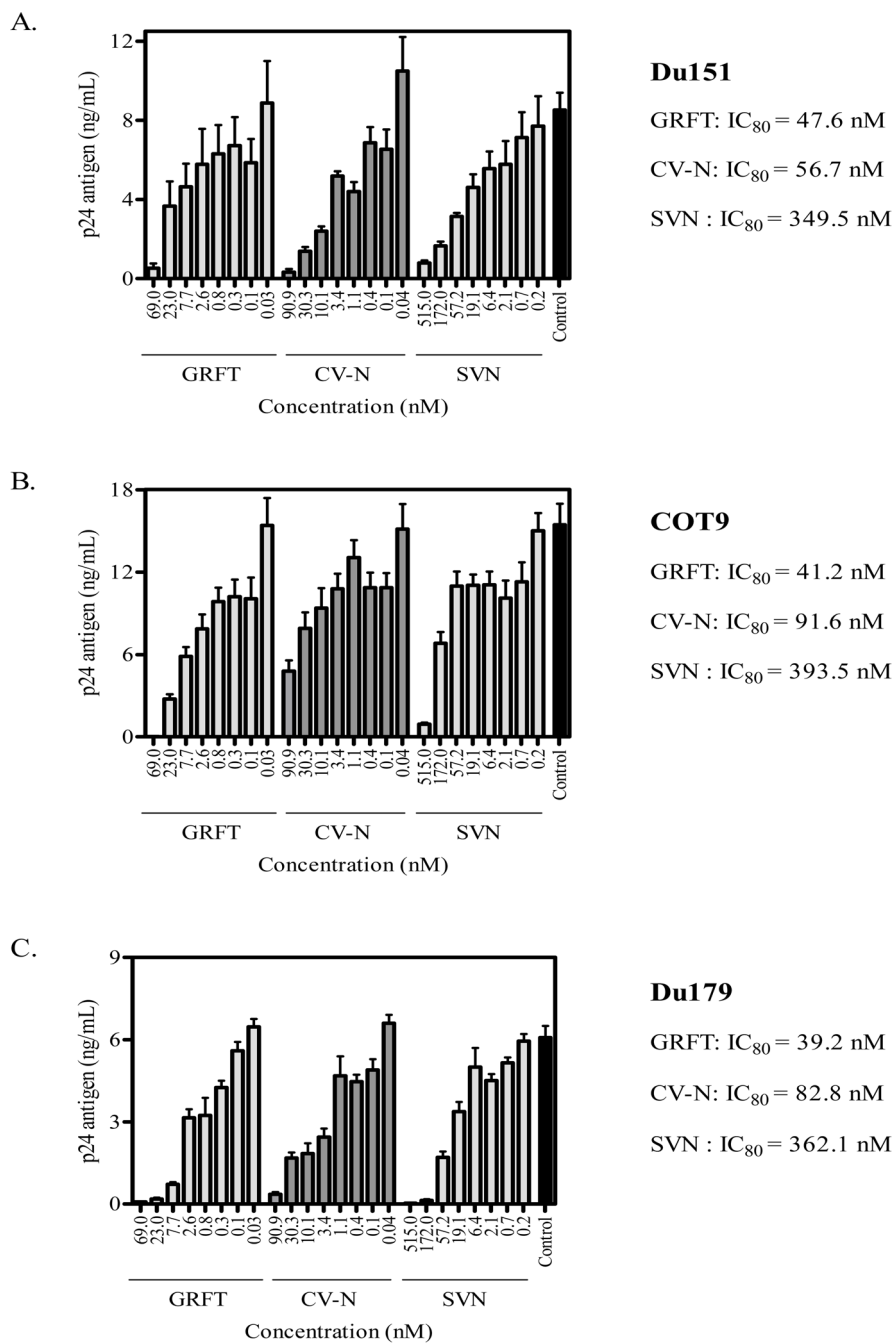


Figure 1. GRFT, CV-N, and SVN inhibit HIV-1 subtype C infection of PBMC
 Three primary isolates, Du151 (A), COT9 (B) and Du179 (C) were treated with increasing concentrations of GRFT, CV-N and SVN before infection of PBMC. Data are shown as the average plus standard deviations of three independent experiments. Untreated virus is shown in black (positive control). The IC₈₀ values of the lectins are indicated next to each graph.

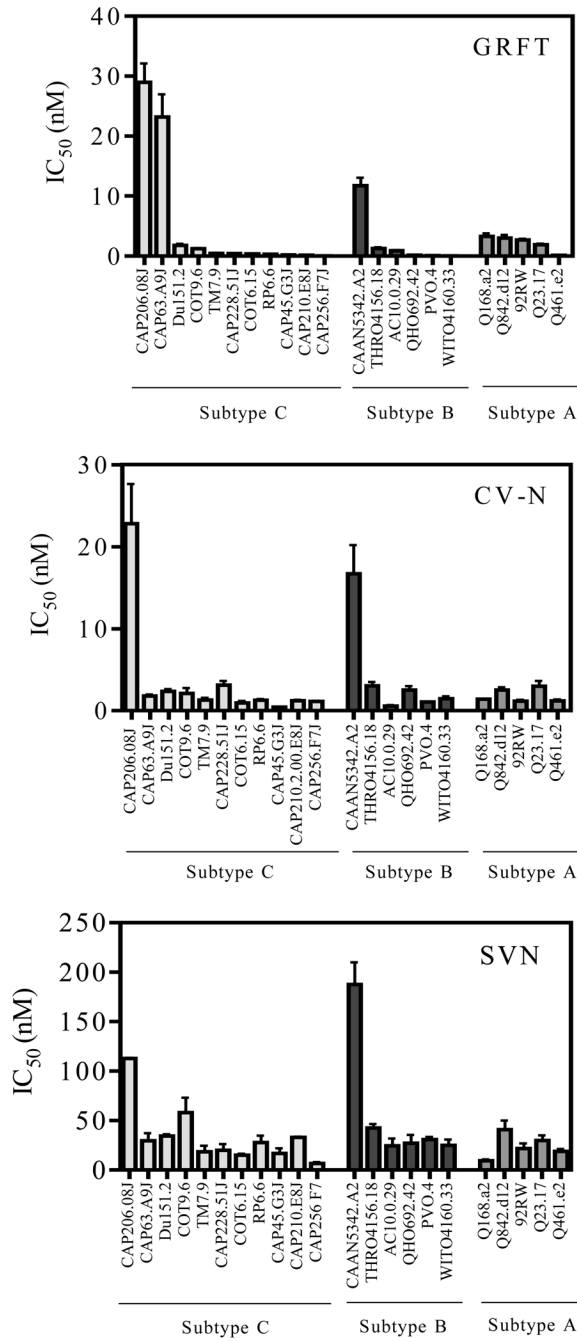


Figure 2. GRFT, CV-N, and SVN inhibit HIV-1 infection in the TZM-bl assay
 Neutralization of HIV-1 subtype C, B and A pseudoviruses by GRFT (A), CV-N (B), and SVN (C). HIV-1 subtype B and A pseudoviruses were used for comparison to subtype C. Each virus was tested at least three times. Pseudoviruses are ranked by GRFT sensitivity.

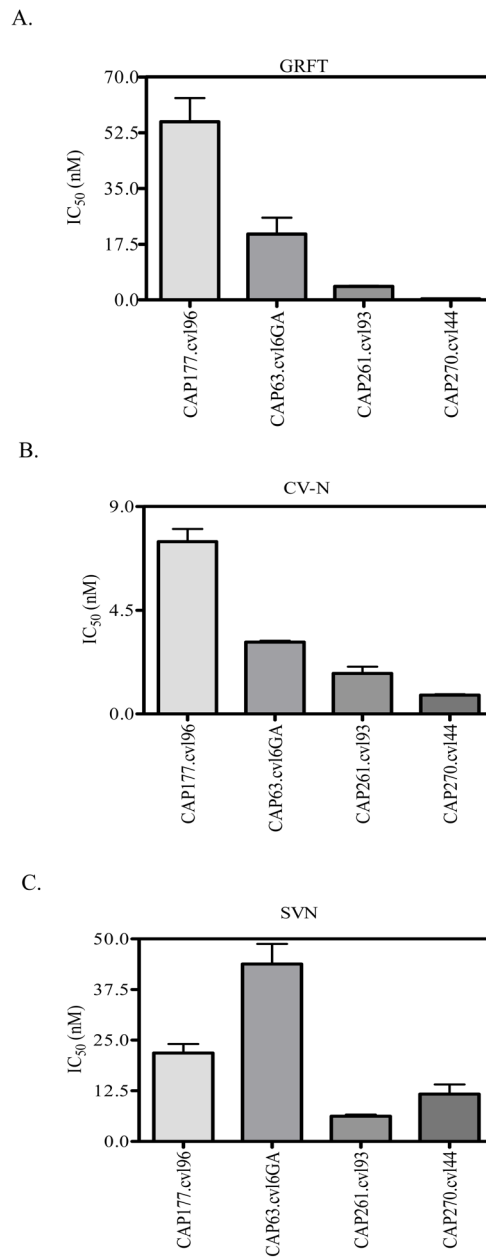


Figure 3. GRFT, CV-N, and SVN inhibit HIV-1 isolates from CVL

Sensitivity of 4 HIV-1 subtype C pseudoviruses containing functional envelope genes amplified from cervico-vaginal lavages against GRFT (A), CV-N (B) and SVN (C) are shown.

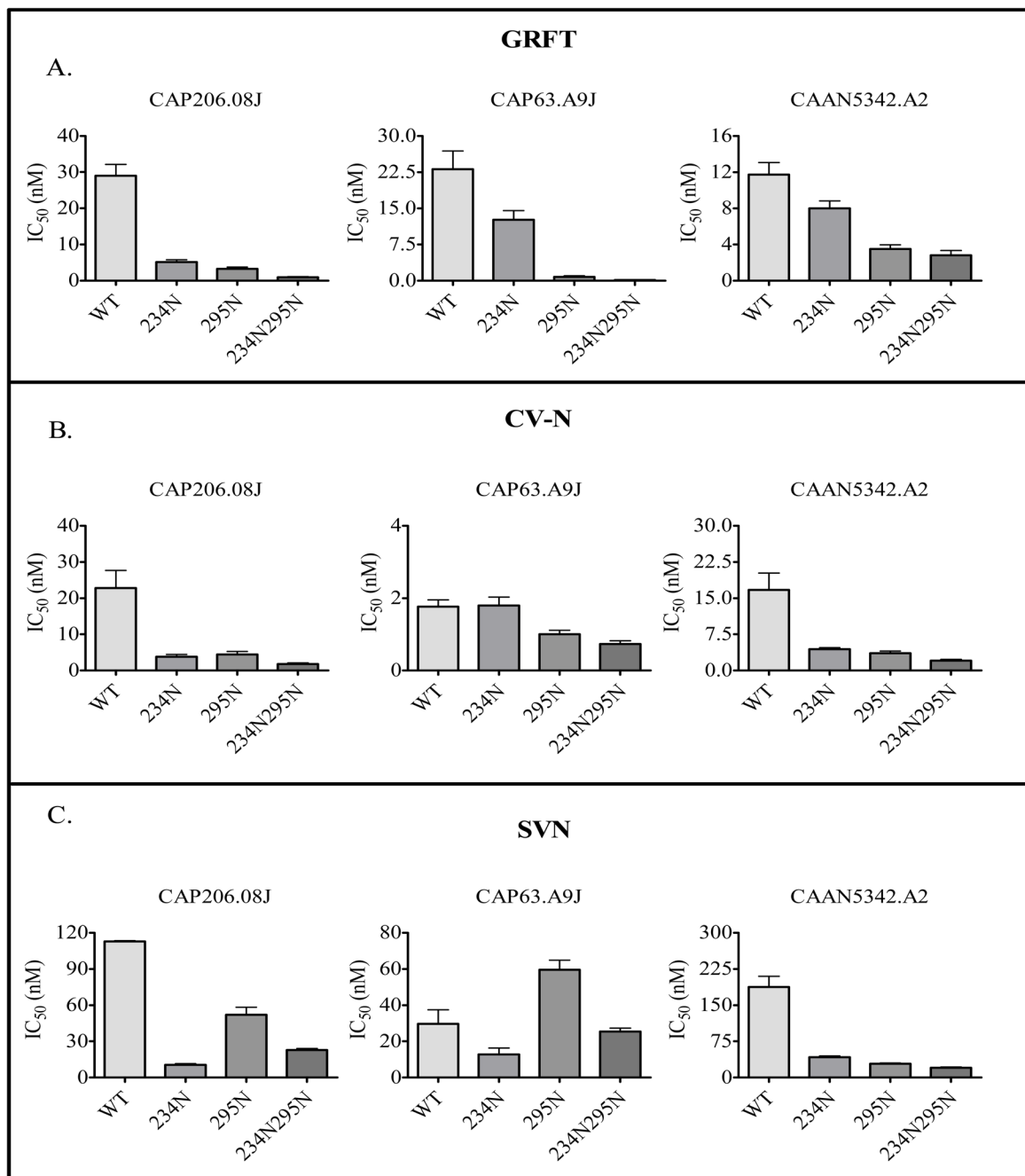


Figure 4. Glycans at positions 234 and 295 increase HIV-1 sensitivity to GRFT, CV-N and SVN
Glycosylation sites at 234 and/or 295 were introduced in CAP206.08J, CAP63.A9J and CAAN5342.A2 by site-directed mutagenesis. Mutant viruses were tested in neutralization assays against GRFT (A), CV-N (B) and SVN (C) in JC53bl-13 cells. Each virus was tested at least three times.

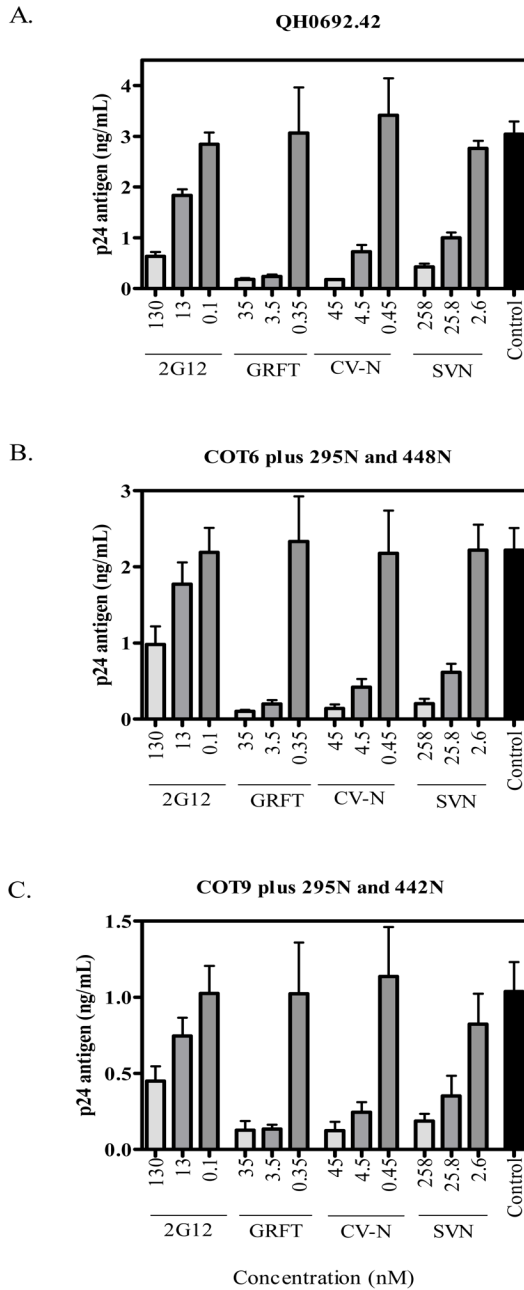


Figure 5. GRFT, CV-N, and SVN compete with the 2G12 mAb for binding to HIV-1 Pseudoviruses QH0692.42 (A), COT6.15-V295N/S448N (B) and COT9.6-V295N/K442N (C), the latter two with a reconstituted 2G12 epitope, were incubated with GRFT, CV-N and SVN prior to capture with the 2G12 mAb. 2G12 was competed against itself as the experimental control. Experiments were done at least 3 times.

Table 1

Pattern of predicted mannose-rich glycans on gp120 of HIV-1 and sensitivity to GRFT, CV-N, and SVN

HIV-1 envelope pseudoviruses	Predicted N-linked mannose-rich glycosylation sites ^a												IC ₅₀ (nM) ^b	
	230	234	241	262	289	295*	332*	339*	386*	392*	448*	GRFT	CV-N	SVN
Subtype C (n = 15)														
CAP177.cv196	x	x	x	x	x	x	x	x	x	x	x	55.9±5.0	7.5±1.1	21.8±5.4
CAP206.08J	x	x	x	x	x	x	x	x	x	x	x	30.9±6.2	18.0±1.2	112.8±1.1
CAP63.A9J	x	x	x	x	x	x	x	x	x	x	x	23.2±6.5	1.8±0.3	29.8±13.3
CAP63.cv16GA	x	x	x	x	x	x	x	x	x	x	x	25.7±2.6	3.1±0.1	43.8±8.5
CAP261.cv193	x	x	x	x	x	x	x	x	x	x	x	4.2±0.4	1.8±0.6	6.2±0.9
Du151.2	x	x	x	x	x	x	x	x	x	x	x	1.5±0.8	2.0±1.0	27.5±9.6
COT9.6	x	x	x	x	x	x	x	x	x	x	x	1.2±0.2	3.9±2.7	57.6±26.9
COT6.15	x	x	x	x	x	x	x	x	x	x	x	0.4±0.2	0.9±0.5	10.8±6.9
CAP228.51J	x	x	x	x	x	x	x	x	x	x	x	0.4±0.1	3.1±0.9	20.1±11.1
CAP270.cv144	x	x	x	x	x	x	x	x	x	x	x	0.3±0.1	0.8±0.07	11.7±4.8
RP6.6	x	x	x	x	x	x	x	x	x	x	x	0.2±0.1	1.1±0.1	11.3±3.3
TM7.9	x	x	x	x	x	x	x	x	x	x	x	0.2±0.1	1.1±0.4	12.1±1.6
CAP45.G3J	x	x	x	x	x	x	x	x	x	x	x	0.2±0.1	0.4±0.1	16.9±8.8
CAP210.E8J	x	x	x	x	x	x	x	x	x	x	x	0.2±0.0	1.2±0.2	33.1±1.1
CAP256.F7J	x	x	x	x	x	x	x	x	x	x	x	0.1±0.0	1.1±0.1	6.7±2.3
<i>Median</i>												<i>0.4</i>	<i>1.8</i>	<i>20.1</i>
Subtype B (n = 6)														

HIV-1 envelope pseudoviruses	Predicted N-linked mannose-rich glycosylation sites ^a											IC ₅₀ (nM) ^b		
	230	234	241	262	289	295*	332*	339*	386*	392*	448*	GRFT	CV-N	SVN
CAAN5342.A2	x				x			x				11.7±2.3	16.7±6.0	187.6±38.6
THRO4156.18				x		x						1.2±0.3	3.0±0.8	42.6±6.9
AC10.0.29	x				x			x				0.9±0.2	0.3±0.1	24.5±12.9
QHO692.42				x					x			0.2±0.1	2.5±0.8	27.3±14.1
PVO.4	x								x			0.1±0.0	1.0±0.1	25.2±14.0
WITTO4160.33		x			x							0.05±0.0	1.5±0.5	24.8±10.1
<i>Median</i>												<i>0.6</i>	<i>2.0</i>	<i>26.2</i>
Subtype A (n = 5)														
Q168.a2	x				x		x	x		x		3.3±0.9	1.4±0.1	9.4±2.3
Q842.d12	x	x			x			x				3.0±0.9	2.5±0.6	40.9±15.8
Q23.17	x				x			x				1.9±0.3	3.0±1.1	29.8±9.0
92RW009	x				x				x			1.8±0.4	1.2±0.3	21.8±8.9
Q461.e2	x				x		x	x		x		0.12±0.06	1.2±0.3	18.9±4.2
<i>Median</i>												<i>1.9</i>	<i>1.4</i>	<i>21.8</i>

^aMannose-rich glycosylations were identified from the amino acid sequence of each envelope clone (Kwong et al., 1998; Leonard et al., 1990). Missing mannose-rich glycosylation sites are marked by X. Viruses are ranked according to GRFT sensitivity.

^bThe IC₅₀ is the concentration of GRFT, CV-N, and SVN that reduced HIV-1 infection by 50%. Mean ± SD of 3 independent is shown.

* Glycans involved in binding the monoclonal antibody 2G12

Table 2

Effect of the 295 glycan on sensitivity to GRFT, CV-N and SVN

Envelope		GRFT	CV-N	SVN
		IC ₅₀ (nM)*		
Du151.2	Wild-type	1.5±0.8	2.0±1.0	27.5±9.6
	V295N	0.03±0.02	1.2±0.4	32.5±4.9
COT9.6	Wild-type	1.2±0.2	3.9±2.7	57.6±26.9
	V295N	0.06±0.04	4.4±3.9	63.0±35.9
COT6.15	Wild-type	0.4±0.2	0.9±0.5	10.8±6.8
	V295N	0.06±0.0	1.1±0.6	16.1±9.2

* IC₅₀ is the concentration of GRFT, CV-N, and SVN that reduces viral infection by 50%.

Cases where the mutant IC₅₀ was more than 5-fold lower than the wild-type virus IC₅₀ are bolded.