Functional characteristics of HIV-1 subtype C compatible with increased heterosexual transmissibility

Brandon L. Walter\textsuperscript{a}, Andrew E. Armitage\textsuperscript{b}, Stephen C. Graham\textsuperscript{c}, Tulio de Oliveira\textsuperscript{d}, Peter Skinhøj\textsuperscript{e}, E. Yvonne Jones\textsuperscript{c}, David I. Stuart\textsuperscript{c}, Andrew J. McMichael\textsuperscript{b}, Bruce Chesebro\textsuperscript{a} and Astrid K.N. Iversen\textsuperscript{b}

\textbf{Background:} Despite the existence of over 50 subtypes and circulating recombinant forms of HIV-1, subtype C dominates the heterosexual pandemic causing approximately 56\% of all infections.

\textbf{Objective:} To evaluate whether viral genetic factors may contribute to the observed subtype-C predominance.

\textbf{Methods:} Chimeric viruses were generated using V1–V3 envelope fragments from a subtype-A/C dually infected woman with preferential genital replication of subtype C. Viral adaptation, spread and cell fusion ability were evaluated \emph{in vitro} using peripheral blood mononuclear cells and HeLa-CD4-CCR5 cell lines, sequencing and cloning. Structural modeling was performed using a crystal structure of gp120-CD4-X5. Phylogenetic analysis was done using subtype-A, subtype-B and subtype-C sequences from blood and cervix of 37 infected women and database sequences.

\textbf{Results:} We identified two envelope motifs, compact V1–V2 loops and V3-316T, which are found at high frequency throughout subtype-C evolution and affect gp120 interactions with CD4 and CCR5, respectively. When a V1-Δ5 deletion or V3-A316T was incorporated into subtype A, each increased viral fusion and spread several fold in peripheral blood mononuclear cell and cell lines with low CCR5 expression. Structural modeling suggested the formation of an additional hydrogen bond between V3 and CCR5. Moreover, we found preferential selection of HIV with 316T and/or extremely short V1–V2 loops in cervix of three women infected with subtypes A/C, B or C.

\textbf{Conclusion:} As CD4\(^{+}\)-CCR5\(^{-}\)-T cells are key targets for genital HIV infection and cervical selection can favor compact V1–V2 loops and 316T, which increase viral infectivity, we propose that these conserved subtype-C motifs may contribute to transmission and spread of this subtype.

\\(©\) 2009 Wolters Kluwer Health | Lippincott Williams & Wilkins

\textit{AIDS} 2009, 23:1047–1057

\textbf{Keywords:} CCR5, cervix, evolution, HIV-1, phylogeny, subtype A, subtype C, transmission

---

\textsuperscript{a}Laboratory of Persistent Viral Diseases, Rocky Mountain Laboratories, National Institute of Allergy and Infectious Diseases, National Institutes of Health, Hamilton, Montana, USA, \textsuperscript{b}MRC Human Immunology Unit, Weatherall Institute of Molecular Medicine, University of Oxford, John Radcliffe Hospital, \textsuperscript{c}Division of Structural Biology, Wellcome Trust Centre for Human Genetics, Oxford, UK, \textsuperscript{d}HRC Pathogen Bioinformatics Unit, South African National Bioinformatics Institute, University of the Western Cape, Cape Town, South Africa, and \textsuperscript{e}Department of Infectious Diseases, Rigshospitalet, Copenhagen, Denmark.

Correspondence to Astrid K.N. Iversen, MRC HLIU, WIMM, University of Oxford, John Radcliffe Hospital, Oxford OX3 9DS, UK.

Tel: +44 0 1865 222498; fax: +44 0 1865 222502; e-mail: astrid.iversen@imm.ox.ac.uk

Received: 7 October 2008; revised: 13 January 2009; accepted: 2 February 2009.

DOI:10.1097/QAD.0b013e32832a1806
Introduction

The current HIV pandemic is dominated by subtype C, which accounts for approximately 56% of the estimated 33 million infections worldwide, despite the concurrent existence of over 50 other subtypes and circulating recombinant forms (CRFs) [1]. First identified in Ethiopia, subtype C has spread through southern and eastern Africa during the last two decades, invading areas formerly dominated by subtypes A and D and is currently causing devastating epidemics in parts of sub-Saharan Africa, Brazil, India and China [2–5].

These transmissions are largely through heterosexual contact or from mother to child. Although founder effects probably can explain the more recent subtype-C epidemics in India, Brazil and China, subtyping of early HIV sequences from southern and eastern Africa strongly suggest that subtype C was introduced into these areas subsequent to subtypes A and D (www.hiv.lanl.gov). As no epidemiological studies have shown any subtype-related differences between heterosexual transmission networks, it has been suggested that subtype-C virus may have a transmission advantage over subtypes A and D [6–8]. Moreover, epidemiological studies [9,10] have demonstrated differences between these subtypes with respect to vaginal shedding and in-utero transmission, subtype C associating with increased risks of both. However, no functional mechanism explaining these differences has so far been identified.

Although many studies have successfully focused on identifying human genetic factors influencing HIV-1 disease progression, these seem to account for just around 10% of the variability in disease progression rates and only some have been demonstrated to affect transmission [11]. Viral genetic variation is also likely to influence pathogenesis and transmission, and short V1–V4 loops, and more neutralization-sensitive viruses, have been suggested to be selected during, or soon after, many sexual transmissions caused by subtypes A and C, but not B [12–16].

Here we asked whether conserved, subtype-specific sequence motifs in subtype C could affect viral replication and transmission. Our results highlight the functional importance of two common subtype-C motifs, short V1–V2 loops and V3-316T, and demonstrate that these are found at high frequency throughout subtype C evolution and may be specifically selected in the genital tract of infected women.

Methods

Study participants

HIV-1-infected women were recruited at Rigshospitalet, Denmark, between 1995 and 1996. Most were untreated; however, some had received monotherapy [azidothymidine (AZT) (8/37), didanosine (ddI) (1/37)] or dual therapy [AZT and ddI (2/37)] for over 6 months [17,18]. The study was approved by the Danish Board of Medical Ethics and patients gave informed consent [17].

Limiting dilution, nested PCR and sequencing

DNA was extracted from peripheral blood mononuclear cell (PBMC) and cervical cell pellets using the QIAamp Blood kit (Qiagen, Valencia, California, USA). V1–V3, C2–C4 and gp160 envelope fragments were amplified by limiting-dilution nested PCR using Advantage 2 Polymerase mix (Clontech, Mountain View, California, USA) and sequenced (sequences are available upon request) [19].

Construction of recombinant HIV-1 clones

Infectious HIV-A and HIV-C chimeric clones with V1–V3 env fragments obtained from blood or cervix of a dually infected patient were generated as described previously for HIV clones 81A-4 and 49-5 [20–23].

PCR amplification, sequencing and cloning of PBMC-adapted virus

Virus-containing supernatants were samples throughout the PBMC infection experiment. Viral RNA was purified using the QIAamp Viral RNA Mini Kit (Qiagen, Valencia, California, USA), reverse transcribed and amplified using the SuperScript One-Step RT-PCR Kit for Long Templates (Invitrogen, Paisley, UK), sequenced and cloned. The reconstructed PBMC-adapted chimeric subtype-A clones were named A20-V1Δ5, A2-T146I, A1-A316T and A1-N302K, according to envelope region and type of mutation.

Virus stocks and infection

Proviral HIV constructs were transfected into RC49 HeLa cells and viral stocks were generated for PBMC, HeLa cell and fusion experiments; titers were expressed as focus-forming units (FFUs) per ml [24,25]. Different virus dilutions [multiplicity of infection (MOI) 0.004–0.1] were used to infect PBMC and cultures were continued for 16 days; p24 levels were measured using an antigen capture assay (HIV-1 p24EIA) as recommended (AIDS Vaccine Program, National Cancer Institute (NCI)–Frederick Cancer Research and Development Center). Infected HeLa cell monolayers were stained for p24 and foci counted as described previously [24,25].

Fusion experiments and immunohistochemical detection

293T cells, which do not express CD4 or CCR5, were transfected with subtype-A, subtype-C or adapted-A chimeric HIV constructs and relative envelope expression levels compared using pooled antisera from HIV-infected patients and fluorescently labeled anti-human IgG (Invitrogen). Data were acquired on a FACSCalibur flow cytometer (Becton Dickinson, Franklin Lakes, New Jersey).
We previously reported preferential genital replication of HIV subtype C in an asymptomatic, untreated woman dually infected by subtypes A and C in blood; her blood-

**Sequence analysis**

Phylogenetic trees were constructed by maximum likelihood using PAUP* 4.0b10 (Sinauer Ltd., Sunderland, Massachusetts, USA) and phyML software (www.atgc-lirmm.fr/phyml/) [26,27]. Nonparametric bootstrap support estimates were generated using 1000 replicates under neighbor-joining tree algorithm using the maximum likelihood substitution model. Tree figures were produced using FigTree1.1 (tree.bio.ed.ac.uk/software/figtree/). Distance analysis was done using MEGA and Kimura-2 parameters; standard errors were estimated using 100 bootstrap runs [28]. The variation analyses were computed using a counting algorithm (http://bioafrica.mrc.ac.za/ JoeTool/counting.html). Frequencies of 316A and T were calculated using 1671 subtype-A and 4165 subtype-C sequences. To decrease the risk of sampling bias in these database-derived sequences, we used only one sequence from each patient or transmission cluster from as many countries as possible (i.e. multiple sequences from known transmission clusters such as wife/husband or mother/baby were excluded). Sequences with translational problems (i.e. frameshift or stop codons) were furthermore excluded as these may either represent nonfunctional virus or could indicate poor sequence quality.

**Molecular modeling and disorder analysis**
The A316T and N302K single amino acid point mutations were modeled on the crystal structure of gp120-CD4-X5 [29] (PDB ID 2B4C) by manually substituting the individual side chains using Coot [30], ensuring by visual inspection that the resulting model maintained good local stereochemistry (Supplementary Fig. S1). The increase in local positive charge at the region of the N302K mutation was confirmed by mapping the molecular electrostatic potential onto the solvent-accessible surface of gp120 (PDB ID 2B4C) and the gp120 N302K model (not shown) using the Adaptive Poisson–Boltzmann Solver (APBS) [31] plug-in for PyMOL (www.pymol.sourceforge.net/; DeLano Scientific) with default parameters. The propensity of gp120 residues to be disordered was predicted from the sequences of primary isolates and adapted strains using the neural-network pattern recognition program regional order neural network (RONN) [32] with default parameters. Molecular graphics were prepared using PyMOL (DeLano Scientific).

**Results**

**Generation of infectious HIV molecular clones**

We previously reported preferential genital replication of HIV subtype C in an asymptomatic, untreated woman dually infected by subtypes A and C in blood; her blood-

We hypothesized that preferential cervical subtype-C infection was associated with viral genetic traits favoring genital infection. To compare the biological characteristics of HIV sequences found in blood and cervix, we amplified and sequenced single-copy V1–V3 envelope fragments from these tissues, as this region contains key determinants of cell tropism, infectivity and chemokine sensitivity [34–36] (Fig. 1a). Sequences were inserted into the pNL4-3ΔV1–V3 vector to generate infectious molecular clones (subtype C: C57, C64, C66, C68, C69 and C85; subtype A: A1, A2 and A20). Infection studies using HeLa cells expressing CD4 and CXCR4 plus or minus CCR5 showed that all virus clones required the CCR5 coreceptor for infection. Furthermore, all viruses were able to infect and spread slowly in monocyte-derived macrophage cultures (data not shown).

Bioinformatic analysis demonstrated that these sequences were common subtype representatives as they clustered within their respective subtypes on phylogenetic trees, and the distance and amino acid variation between them and other subtype-A and subtype-C sequences were comparable to the medium intrasubtype distance and variation (supplementary Table S1, data not shown). Thus, the viral sequences from our patient may provide information on how subtypes A and C compare functionally in vivo.

**In-vitro PBMC replication and cell fusion**

In PBMC cultures, we noted marked subtype-specific differences in live virus production. Whereas subtype-C titers peaked at 4–6 days postinfection, subtype-A titers rose more slowly over the course of 16 days (Fig. 1b). Throughout the PBMC infection experiment, viral supernatants were removed and used to infect JC37 cells, which expressed low levels of CCR5 and high levels of CD4. Regardless of sampling time, supernatants from the subtype-C infections produced large foci (10–20 nuclei/focus; Fig. 1d). In contrast, supernatants from the subtype-A infections at days 4–8 generated mostly small foci (1–2 nuclei/focus). However, subtype-A foci increased markedly in size at days 12–16, coincident with increased replication in PBMC, suggesting that adaptation occurred during in-vitro culture of these viruses.

**Fusogenicity studies**

We examined the ability of the viral envelope protein to fuse with cell lines expressing CD4 and CCR5 by transfecting 293T (CD4-/-, CCR5-/-) cells with plasmid DNA of subtype-A and subtype-C infectious clones. The transfected cells expressed similar cell surface viral envelope levels by flow cytometry analysis at 48 h posttransfection. When these cells were overlaid for
24 h on RC49 cells (CD4<sup>low</sup>, CCR5<sup>high</sup>), both subtypes A and C induced a high level of fusion (Fig. 1e). By contrast, when they were overlaid onto JC37 cells (CD4<sup>high</sup>, CCR5<sup>low</sup>), subtype-A viruses generated small HIV-positive fusion foci ranging in size from 1 to 9 nuclei (mean 2 nuclei per focus), whereas subtype-C foci contained from 1 to more than 100 nuclei per focus (mean 20 nuclei). Thus, HeLa cells with low CCR5 levels were fused more efficiently by subtype-C viruses than by subtype A viruses, but both subtypes resulted in extensive fusion on cells expressing high CCR5 levels.

**Selection of adapted subtype-A mutants in PBMC**

During replication in PBMC (Fig. 1b), subtype-A viruses attained the high titers seen early in subtype C infection only slowly, and the early and late supernatants from subtype A infections differed in their ability to fuse JC37 cells, suggesting that mutant subtype A viruses had been selectively amplified during in-vitro culture. To search for evidence of adaptation, we reverse transcribed, amplified and sequenced viral subtype-A RNA from day 16 and found four distinct envelope sequence changes targeting the V1 and V3 loops. Using these sequences, we
generated new chimeric viruses, A1–A316T, A1–N302K, A2–T146I and A20–V1Δ5, which, apart from the indicated changes, were otherwise identical to their parental virus (Fig. 1a). These adapted viruses displayed a distinct functional phenotype, which differed from their parental subtype-A clones but was similar to the chimeric subtype-C clones, that is, they spread rapidly in PBMC and produced large foci in JC37 cells (Fig. 1c,f), cell types that both express low levels of CCR5.

The influence of CCR5 expression levels on virus titers
To determine whether CCR5 expression levels influenced viral replication in CD4+, CCR5+ HeLa cells, subtype-A, subtype-C and PBMC-adapted subtype-A clones were used to infect JC37 and RC49 cells. In RC49 (CD4low, CCR5high) cells, all viruses gave high titers (2.5 × 10^5 – 3.7 × 10^6 FFU/ml) at 5 days postinfection (Table 1). In contrast, in JC37 (CD4high, CCR5low) cells, all three subtype-A clones produced low-virus titers (1.9 × 10^3 – 1.7 × 10^4 FFU/ml), whereas subtype-C and all four PBMC-adapted subtype-A clones produced high titers similar to those seen on RC49 cells.

Structural modeling of envelope variation
Structural modeling was performed to help understand the molecular consequences of the four subtype-A in vitro adaptations (Supplementary Fig. S1). The V1–V2 envelope region lies close to the CD4-binding site, whereas the V3 region interacts with two parts of CCR5 (Supplementary Fig. S1a). The V3-A316T substitution may strengthen HIV-CCR5 interactions as the additional polar group is in an appropriate position to form a new hydrogen bond to CCR5, most likely independently of the surrounding genomic context (Supplementary Fig. S1b). Similarly, the N302K substitution adds another positive charge to the V3 stem, presumably strengthening CCR5 binding, as a crystal structure of V3 and the N-terminal negatively charged CCR5 tail has shown it to interact directly with position 302 [37].

The interactions between the CD4 receptor and the V1–V2 envelope region are less precisely understood as no high-resolution structures of V1–V2-containing gp120 envelopes exist. The loss of a V1-glycosylation site in A2–T146I and the A20–V1Δ5 deletion might enhance CD4 receptor contact, perhaps by decreasing steric hindrance. We used a neural-network technique to analyze the propensity for disorder of the envelope proteins, as disordered regions can combine high-binding specificity with modest affinity [32,38] (Supplementary Fig. S1c). We found significant differences in disorder probability between subtypes A, A20–V1Δ5 and C in the V1–V2 region, suggesting variation in the way these sequences might bind to CD4 and/or conformationally mask the CD4-binding site; however, both wild-type-A and wild-type-C chimeric viruses utilized the CD4 receptor efficiently in our functional assay.

Analysis of in-vivo occurrence of mutations selected in vitro
We examined the frequency of the four subtype-A in vitro adaptations in natural HIV infections. Two substitutions were found occasionally in natural infections in some subtypes, however, rarely (146I, 2%) or never (302K) in subtype A (Table 2). By contrast, V3–316T was seen in 68% of all subtype-C sequences, but at much lower frequencies, if at all, in the other subtypes (25–0%) (Table 2, Fig. 2). Furthermore, the V1–Δ5 subtype-A deletion, which shortens the loop, resembled the short V1–V2 loops typically found in subtype C but not in subtypes A, B and D (Table 2). The high frequency of V3–316T and short V1–V2 loops in subtype C, which both increase HIV spread in vivo, suggest these motifs may affect HIV replication in infected patients.

Phylogenetic analysis of 316T selection
While deletions cannot be incorporated in phylogenetic analysis at present, we could examine V3–316T selection. We generated a phylogenetic tree based on subtype-C sequences from 1985 to 2005 and superimposed the nature of the amino acid at position 316 (Fig. 2a, Supplementary Fig. S2). The earliest sequences are from the Democratic Republic of Congo (DRC) where HIV is thought to have evolved and diversified. The phylogenetic tree demonstrates that both 316A and 316T could be found in the DRC sequences at the base of the tree. However, the deep lineages and branches of the tree mostly carry 316T, suggesting that this residue has been generally advantageous to subtype-C spread. Whereas 316V can be found sporadically, alanine seems to spread locally, especially in more recent sequences from South Africa and Botswana. We studied the distribution of 316A and 316T in larger alignments of shorter envelope sequences dating from 1990 to 2005 to see if this observation could be substantiated (Fig. 2b). We found that the frequency of 316T indeed had decreased from around 80% in 1990 to approximately 68% in 2005. This suggests that there may be alternate selective advantages for 316A in some situations in vivo and/or that other

---

**Table 1. Comparison of chimeric virus infectivity titers in supernatant fluid from infected JC37 and RC49 cells.**

<table>
<thead>
<tr>
<th>Clone</th>
<th>Titer JC37 CCR5low</th>
<th>Titer RC49 CCR5high</th>
<th>Ratio (RC49/JC37)</th>
</tr>
</thead>
<tbody>
<tr>
<td>A1</td>
<td>9.7 × 10^4</td>
<td>1.0 × 10^5</td>
<td>10.31</td>
</tr>
<tr>
<td>A2</td>
<td>1.7 × 10^4</td>
<td>1.6 × 10^5</td>
<td>9.41</td>
</tr>
<tr>
<td>A20</td>
<td>1.9 × 10^4</td>
<td>2.5 × 10^4</td>
<td>131.58</td>
</tr>
<tr>
<td>C66</td>
<td>1.6 × 10^5</td>
<td>3.7 × 10^4</td>
<td>2.31</td>
</tr>
<tr>
<td>A1-V3-A316T</td>
<td>6.9 × 10^4</td>
<td>6.7 × 10^4</td>
<td>0.97</td>
</tr>
<tr>
<td>A1-V3-N302K</td>
<td>4.2 × 10^4</td>
<td>5.4 × 10^4</td>
<td>1.29</td>
</tr>
<tr>
<td>A2-V1-T146I</td>
<td>7.9 × 10^4</td>
<td>6.2 × 10^4</td>
<td>0.78</td>
</tr>
<tr>
<td>A20-V1Δ5</td>
<td>9.3 × 10^4</td>
<td>9.9 × 10^4</td>
<td>1.06</td>
</tr>
</tbody>
</table>

FFU, focus-forming unit.
changes sometimes are sufficient to favor transmission of subtype C.

**Analysis of HIV-316T distribution in blood and cervix**

To analyze the prevalence of 316T in blood and cervix, we obtained samples from 36 additional women infected by HIV subtypes A (n = 3), B (n = 28) or C (n = 5). Whereas single-copy envelope sequences could be amplified from PBMC from all women, no HIV was found in the cervices of four (19).

Most subtype-B infected women carried virus with V3-316A, but four had either 316T, 316T/M, 316S or 316V virus in both samples (Fig. 3a). Moreover, we found V3-316 compartmentalization between blood and cervix in one subtype-B-infected woman (1/28, 4%); all genital viruses carried 316T (5/5), whereas only 316A could be found in PBMC (7/7) (Fisher’s exact test, P < 0.001; Fig. 3b).

Of the five subtype-C-infected women, only one carried HIV-316A, and no virus could be amplified from her genital sample; the rest had either 316T or 316T/M in both PBMC and cervix. In one of these patients with HIV-316T, we were able to analyze gp160 from vagina as well as cervix and PBMC (Fig. 3c). Although 28% of the cervical sequences had very short V1–V2 loops (55 amino acids), such short loops were not found in blood; most vaginal sequences had intermediate V1–V2 loops. Overall, V1–V2 loop lengths differed significantly between genitalia and blood when grouped into either short (≤69 amino acids) or long (>69 amino acids) loops (Fisher’s exact test, P = 0.02).

While our results demonstrate that subtype-A and subtype-B virus with 316A can replicate in the cervix, they also highlight that selective forces in the genital tract may favor replication of virus with 316T even in subtype-B-infected women. This is reminiscent of the preferential cervical replication of subtype-C virus in our dually infected woman. Furthermore, we demonstrate that in a woman carrying only HIV-316T, selective forces in the genital tract may favor shorter V1–V2 loops than selection in blood.

**Discussion**

Subtype C is responsible for an ever-increasing fraction of new HIV infections worldwide, especially in areas where heterosexual transmission predominates, and it now prevails even in regions where subtype A or D or both used to be common (1,2,39,40). This suggests that there are biological factors favoring transmission of subtype C, but no plausible mechanism has been suggested.

Our study of infectious, chimeric HIV clones carrying V1–V3 sequences from a woman dually infected with subtypes A and C, but with preferential cervical subtype-C replication, combined with the analysis of HIV sequences from blood and cervix from 36 additional women, lead to four findings suggesting that subtype C generally has a relative replication advantage compared with most subtype-A and subtype-B viruses in the female genital tract.

First, we identify a V3-envelope signature sequence (316T) at high frequency in subtype C, which might strengthen the interaction between HIV and CCR5, the main coreceptor in the female genital tract (41–45). We find that the effect of 316T may be independent of the surrounding genomic context and show the effect of an A316T substitution in subtype A. Second, we find that
Fig. 2. The presence of V3-316T and V3-316A in subtype A and C over time and frequency in subgroup M. (a) Phylogenetic tree of a 219 base pair env region (HXB2 7050-7271) from 163 subtype-C sequences (1985–2005), including our patient sequences. The scale bar shows the branch length equal to five nucleotide changes per 100 bases (0.05). Sequences carrying 316 A, T and V are blue, red and purple respectively; branches where all sequences have the same amino acid at position 316 are collapsed and shown as colored triangles (uncollapsed tree with GenBank accession numbers, Supplementary Fig. S2). Ancestral branches for India and South America are labeled with /C3 and /C3/C3, respectively. (b) Graphs show the prevalence of 316A and T in subtype A (blue) and C (red), respectively, from 1990 to 2005 (number of sequences, subtype A/C: 1985–1990, n = 82/261; 1991–1995, n = 147/356; 1996–2000, n = 614/1519; 2001–2005, n = 166/993). (c) Cladogram of 263 V3 env sequences showing the distribution of 316A (blue) and T (red) in all subgroup M subtypes (M = subgroup M root). AR, Argentina; BR, Brazil; BW, Botswana; DRC, Democratic Republic of Congo; ET, Ethiopia; IL, Israel; IN, India; KE, Kenya; SE, Sweden; TZ, Tanzania; UG, Uganda; UY, Uruguay; ZA, South Africa.
selection for 316T can occur even in the cervix of a minority of untreated subtype-B-infected women. Third, we demonstrate that a subtype-A V1-deletion augments virus production in cell lines expressing low levels of CD4 or CCR5 and propose that the shortening of the V1 loop increases viral interactions with the CD4 receptor. Fourth, we find that genital selection in a subtype-C-infected woman favors outgrowth of very short V1–V2 loops, despite subtype C already having the most compact V1–V2 loops of all HIV subtypes.

We found that subtype C had a greater ability to replicate in PBMC cultures than subtype A and mediated more extensive virus-induced fusion of cells expressing low CCR5 levels. However, we observed marked changes in subtype-A replication patterns over time and identified adaptations in either the V1 (T114I, V1-D5) or V3 (N302K, A316T) region, which are known to interact with the CD4 and CCR5 receptors, respectively. These adaptations enhanced virus replication and fusion comparable to what we found using the wild-type subtype-C clones. Whereas two were found rarely, if at all, in HIV sequences obtained from infected persons, two others (A316T and V1-D5) resembled sequence motifs in the subtype C sequences from our patient and in most subtype C sequences from natural infections. In other subtypes, and in most CRFs, alanine is preferred at position 316, with the notable exception of CRF_02_AG, which carries threonine. Analogous to subtype-C spread in southern and eastern Africa, CRF_02_AG has spread throughout western Africa, primarily via heterosexual contact [1,46].

Both subtype-A and subtype-C chimeric viruses interacted efficiently with the CD4 receptor; however, the V1-D5 adaptation augmented this interaction and increased infectivity several-fold in both cell lines and PBMC. The CD4-binding region is predicted to be shielded by the V1–V2 loop and shorter loops have been associated with increased susceptibility to antibody neutralization [13–15]. As short V1–V4 loops and more neutralization-sensitive HIV variants have been found in
Early subtype-A and subtype-C infections, it was suggested that these were either preferentially transmitted or rapidly selected in the recipient [12–15]. We demonstrate that genital selection pressures in subtype-C-infected women may sometimes favor virus with more compact V1–V2 loops than what is commonly found in blood. Consequently, whether transmission of such variants is preferential or at times simply occurs by chance, remains to be elucidated.

Structural modeling suggested that the A316T effect might be independent of the surrounding genomic background, as a new hydrogen bond may form between the polar threonine and CCR5. The importance of position 316 in gp120-CCR5 interactions is indicated by the induction of drug-resistance mutations at this site by two different CCR5 inhibitors (A316T, maraviroc; A316V, AD101) [47,48]. CCR5-binding studies [47,48] have demonstrated that these polar and nonpolar substitutions crucially affect HIV-CCR5 interactions through different mechanisms.

Our phylogenetic analysis demonstrated that 316T must have been favorable for subtype-C transmission. However, it also revealed that 316A seems to spread locally, especially in recent sequences from South Africa and Botswana. This implies that viruses with 316T incur a viral fitness cost even in subtype C, as has been directly demonstrated for maraviroc-resistant subtype-B virus in subtype C-infected women may sometimes favor virus with more compact V1–V2 loops than what is commonly found in blood. Consequently, whether transmission of such variants is preferential or at times simply occurs by chance, remains to be elucidated.

In conclusion, our study suggests that subtype C has evolved in ways compatible with increased sexual transmissibility. Although other envelope regions, additional viral genes and long terminal repeat promoter differences [55] in combination with epidemiological dynamics, such as founder effects, also must be important factors, these subtype-C motifs may contribute to its current dominance in southern and eastern Africa, which facilitated subsequent worldwide spread.

Acknowledgements

We thank the patients, Andrew Rambaut, Kim Hasenkrug, Rachel Lacasse, Angela Vincent and Nick Willcox for helpful comments; Anita Mora for graphics, Kevin Braughton and Ron Messer for technical assistance.

A.K.N.I. initiated and designed the overall study; A.K.N.I. and B.C. planned and supervised different parts of the project; A.K.N.I., B.C. and B.L.W. analyzed data and wrote the article; B.L.W., A.E.A. and A.K.N.I. performed experiments; S.C.G., E.Y.J. and D.I.S. performed the structural modeling and RONN analyses; T.d.O. and A.K.N.I. performed the phylogenetic analysis; A.K.N.I. and P.S. provided patients, clinical data and reagents; and A.J.M. contributed reagents. All authors discussed the results and approved the final version of the article.

Funding was provided by the Danish AIDS Foundation, the Nuffield Dominions Trust, Cancer Research UK, the
References


Supplementary documentation.

**Supplementary Fig. 1 Structural modeling.**

(a) Proposed gp120/CD4/CCR5 complex. The gp120 V3 loop makes two independent contacts with CCR5: the V3 crown interacts with the extra-cellular surface loops of CCR5, while the stem and base interact with the CCR5 tail. Only a monomer of the presumed trimeric interaction is shown. (b) Structure of HIV gp120 bound to the C-terminal 2 domains of CD4 and the antigen-binding fragment of the neutralizing antibody X5, which targets the CCR5 binding site[27] (PDB-ID 2B4C). Green spheres denote N302 and A316 in the V3 loop stem and crown, respectively. (c) RONN disorder predictions of the V1-V3 region of subtype-C (black), subtype-A (grey) and PBMC-adapted viruses; A2-T144I and A20-V1Δ5 sequences are shown in green while A1-A316T, A1-N302K were indistinct from their parent virus. Values above 0.5 are predictive of disorder[29].

**Supplementary Fig. 2 Uncollapsed phylogenetic tree with GenBank accession numbers.**

**Supplementary Table 1 Bioinformatic analysis.**

<table>
<thead>
<tr>
<th>Average variation</th>
<th>Mean Genetic Distance</th>
</tr>
</thead>
<tbody>
<tr>
<td>pt A</td>
<td>0.013 (SE 0.001)</td>
</tr>
<tr>
<td>subtype A</td>
<td>0.124 (SE 0.006)</td>
</tr>
<tr>
<td>pt/subtype A</td>
<td>0.107 (SE 0.005)</td>
</tr>
<tr>
<td>pt C</td>
<td>0.008 (SE 0.001)</td>
</tr>
<tr>
<td>subtype C</td>
<td>0.121 (SE 0.007)</td>
</tr>
<tr>
<td>pt/subtype C</td>
<td>0.113 (SE 0.006)</td>
</tr>
<tr>
<td>Average distance</td>
<td></td>
</tr>
<tr>
<td>subtype A</td>
<td>0.126 (SE 0.007)</td>
</tr>
<tr>
<td>pt A/subtype A</td>
<td>0.106 (SE 0.011)</td>
</tr>
<tr>
<td>subtype C</td>
<td>0.121 (SE 0.007)</td>
</tr>
<tr>
<td>pt C/subtype C</td>
<td>0.118 (SE 0.010)</td>
</tr>
</tbody>
</table>

Average amino acid variation and genetic distance analyses within and between the patient and database sequences. pt, Patient; SE, standard error.