Identification and Characterization of Conserved and Variable Regions in the Envelope Gene of HTLV-III/LAV, the Retrovirus of AIDS

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Summary
To determine the extent and nature of genetic variation present in independent isolates of HTLV-III/LAV, the nucleotide sequences of the entire envelope gene and parts of gag and pol were determined for two AIDS viruses. The results indicated that variation throughout the viral genome is extensive and that the envelope gene in particular is most highly variable. Within the envelope, changes were most prevalent within the extracellular region where clustered nucleotide substitutions and deletions/insertions were evident. Based on predicted secondary protein structure and hydrophilicity, these hypervariable regions represent potential antigenic sites. In contrast to the hypervariable regions, other sequences in the extracellular envelope and the overall envelope structure (including 18 of 18 cysteine residues), as well as most of the transmembrane region, were highly conserved.

Introduction
The extreme morbidity, high mortality, and epidemic proportions of the acquired immune deficiency syndrome (AIDS) have led to intense scientific efforts to elucidate the disease’s pathogenesis and develop effective preventive and treatment measures. Fundamental to this effort was the identification and characterization of HTLV-III/LAV as the causative agent of AIDS (Barre-Sinoussi et al., 1983; Miescher, 1983, 1984; Schubach et al., 1984; Soragnadharan et al., 1984) and the subsequent molecular characterization of the viral genome (Hahn et al., 1984; Shaw et al., 1984; Alizon et al., 1984; Luzic et al., 1984; Ratner et al., 1985a; Wain-Hobson et al., 1985; Sanchez-Pescador et al., 1985; Muesing et al., 1985).

A troublesome finding, given its potential clinical implications, made early in the genetic analysis of HTLV-III/LAV was that independent virus isolates were heterogeneous in their genomic restriction enzyme cleavage patterns. This observation was extended by electron microscopic heteroduplex studies (Hahn et al., 1985). DNA sequencing (Ratner et al., 1985a; Wain-Hobson et al., 1985; Sanchez-Pescador et al., 1985; Muesing et al., 1985), and additional restriction enzyme mapping (Benn et al., 1985), all of which indicated that substantial genomic variation is present in different AIDS virus isolates and that the envelope might be particularly variable compared to other viral genes (Ratner et al., 1985b; Rabson and Martin, 1985). From these studies, however, it was impossible to estimate accurately the extent and nature of genomic heterogeneity present among independent AIDS virus isolates because restriction mapping and heteroduplexing lacked sufficient sensitivity and because sequence comparisons were performed on only three virus isolates, two of which were relatively closely related to each other.

Because of these limitations, and because a more complete understanding of genomic heterogeneity would undoubtedly be critical to elucidating the virus’s evolutionary history and pathobiology, we undertook a study of genetic variation in five independent HTLV-III/LAV isolates. Here we present an analysis of the nucleotide and deduced amino acid sequences of the complete envelope genes and parts of gag and pol for HTLV-III/LAV isolates obtained from two Haitian patients with AIDS and compare these to the published sequences of HTLV-III (Ratner et al., 1985a), LAV (Wain-Hobson et al., 1985), and ARV (Sanchez-Pescador et al., 1985). The results of these studies provide considerable new insight into the extent and nature of genetic variation present in field isolates of HTLV-III/LAV, and provide important information regarding the rate and mechanisms by which these changes occur in vivo.

Results
Nucleotide Sequence Determination and Analysis
Isolates of HTLV-III/LAV were obtained from peripheral blood mononuclear cells. Isolate RF was obtained in 1983 as previously described (Popovic et al., 1984) and was propagated in an immortalized T cell line from which it was molecularly cloned (Hahn et al., 1985a). Isolate WMJ-1 was similarly transmitted from the peripheral blood mononuclear cells of a Haitian infant with AIDS to an immortalized T cell line (Hahn et al., 1986). Molecular clones of both RF (designated HAT-3) and WMJ-1 were prepared as described in Experimental Procedures.

The genome of HTLV-III/LAV consists of long terminal repeat (LTR) elements; gag, pol, and env genes; two regions of unknown significance designated soF or short open reading frame and 3′ orF for 3′ open reading frame; and a third region previously thought to be noncoding (NCR) but now known to encode the first exon of the HTLV-
Table 1. Sequence Comparison of Five Independent AIDS Virus Isolates

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<thead>
<tr>
<th>Nucleotide Differences Compared to BH-10 (number of nucleotide changes/number of nucleotides sequenced)</th>
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<tr>
<td><strong>Leader Sequence</strong></td>
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<td><strong>HAT-3</strong></td>
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<td><strong>ARV-2</strong></td>
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<td><strong>4.7%</strong></td>
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<td><strong>1.8%</strong></td>
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<td><strong>WMJ-1</strong></td>
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<td><strong>3.3%</strong></td>
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<tr>
<th>Amino Acid Differences Compared to BH-10 (number of amino acid changes/number of amino acids sequenced)</th>
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<tr>
<td><strong>HAT-3</strong></td>
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<td><strong>24/2039</strong></td>
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<td><strong>ARV-2</strong></td>
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<td><strong>ARV-2</strong></td>
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<td><strong>LAV-la</strong></td>
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<td><strong>WMJ-1</strong></td>
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*For WMJ-2, a highly related viral clone from this same patient, the complete sequence of the gag p24 and p15 is shown (Hahn et al., 1986).*

Nucleotide and amino acid sequence comparisons of five independent AIDS virus isolates using the BH-10 clone of HTLV-IIIB as a reference (Ratner et al., 1985a). Deletions and insertions, as illustrated in Figures 1 and 2, are included in the analysis. Data for ARV-2 and LAV-la have been reported elsewhere (Sanchez-Pescador et al., 1985; Wain-Hobson et al., 1985). Data for HAT-3 and WMJ-1 are from the present study. As discussed in the text and shown here, the entire LTR, NCR[tat], envelope, and 3' orf of HAT-3 were sequenced. For WMJ-1, the entire envelope gene was sequenced.

The distribution of sequence differences was not uniform throughout the viral genomes. Instead, when divergent genomes were compared, changes were found to be considerably more prevalent in the envelope and 3' orf than in other genes (Table 1). In the envelope gene in particular, BH-10 differed from HAT-3 in 12.5% of nucleotides (19% amino acids), from ARV-2 in 9.8% of nucleotides (15.7% amino acids), and from WMJ-1 in 9.3% of nucleotides (14.0% amino acids). In the same region, HAT-3 differed from ARV-2 in 14.7% of nucleotides (18.4% amino acids) and from WMJ-1 in 11.3% of nucleotides (16.5% amino acids). ARV-2 and WMJ-1 differed from each other in this region by 10.0% of nucleotides (16.2% amino acids). The most conserved areas in the genomes of HAT-3, LAV-la, BH-10, and ARV-2 were the gag and pol genes, which differed among the viruses in less than 6% of nucleotides and less than 7% of amino acids. In these regions, nucleotide sequence changes were almost exclusively due to point mutations in contrast to env where
Clustered nucleotide changes involving in-frame deletions, insertions, and/or duplications were common. For example, compared to BH-10, there were 48 bp of insertions and 21 bp of deletions in the extracellular envelope of HAT-3 as compared to only 6 bp of insertions and deletions in all of its analyzed gag, pol, and sor sequences. Similarly, there were 33 bp of insertions and 36 bp of deletions in the extracellular envelope of ARV-2 (compared to BH-10) but only a 6 bp insertion and no deletions in all of its gag, pol, and sor. Even in LAV-1a, which is more closely related to BH-10 than are the other sequenced viruses, there was a 15 bp insertion (duplication) in the exterior envelope gene but no insertions or deletions in gag, pol, or sor. Not included in this comparative analysis is a single direct repeat at the gag/pol junction of BH-10 that is not present in any of the other isolates.

Another difference in the types of mutations present in the env gene compared to the gag gene of these viruses was in the proportion of silent third base pair changes. Excluding deletions and insertions, compared to BH-10 the mutations in the external env genes of ARV-2, HAT-3, and WMJ (clone WMJ-2 from patient WMJ; see Hahn et al., 1986) were in the third position of the codon in 36%, 37%, and 34% of instances leading to 32%, 25% and 42% amino acid changes, respectively. Conversely, in the gag genes of these same viruses, 63%, 56%, and 66% of nucleotide changes were in the third place of the codon leading to 10%, 3.5% and 0% amino acid changes, respectively. Thus more than half of the single nucleotide changes in env occurred in the first or second codon position, and even for third position changes, many of these led to amino acid changes. Conversely, only a minority of first or second codon position changes occurred in gag, and in this gene the third position changes were almost uniformly silent. These findings are consistent with two interpretations; first, that structure-function relationships lead to stronger conservation of amino acid sequence in gag than in env, and, second, that non-silent nucleotide changes in env may actually be under positive selection pressure. Whether or not the latter interpretation is correct must await a much more complete understanding of the host immunologic response to HTLV-I/LAV.

Alignment and Comparison of Envelope Sequences

An alignment of the envelope nucleotide and amino acid sequences of HAT-3, WMJ-1, BH-10, LAV-1a, and ARV-2 is shown in Figures 1 and 2. A methionine codon at position 8 of the BH-10 envelope open reading frame most likely marks the beginning of the gene, with a hydrophobic region between position 17 and 37 representing the potential leader sequence. It is known from the amino acid sequence of the mature envelope glycoprotein that this leader sequence, or signal peptide, is cleaved from the envelope precursor protein during envelope maturation (Allen et al., 1965). The ensuing peptide sequence from position 38 to 518 (BH-10 sequence) is slightly hydrophilic and contains numerous potential N-linked glycosylation sites, ranging from 22 in WMJ-1 to 29 in HAT-3. This region corresponds to the major exterior envelope glycoprotein (gp120). A characteristic arginine-rich hydrophobic stretch marks the cleavage site for the processing of the envelope precursor gp160 into the exterior gp120 and membrane-bound gp41 (Allen et al., 1965; DiMarzo-Veronese et al., 1985). The latter protein includes an apparent hydrophobic membrane-spanning segment, a hydrophilic anchor sequence, and an additional carboxy-terminal stretch of 150 residues whose function is presently unknown.

A striking feature of the five envelope sequences was the conservation of cysteine residues (Figure 2). Within the extracellular envelope, each of the 18 cysteine residues was conserved in all five viruses. Within the transmembrane protein, all but one cysteine residue in its extreme carboxy-terminus was conserved. This finding argues for a highly conserved "macrostructure" of different HTLV-III/LAV envolopol glycoproteins. Despite this overall conservative structure, numerous amino acid changes were evident throughout the envelope glycoprotein. In the transmembrane region these were generally the result of isolated nucleotide point mutations leading to single amino acid substitutions. In the extracellular region, changes in amino acid sequence resulted from both nucleotide point mutations and in-frame deletions, insertions, or duplications of blocks of nucleotides. The resulting amino acid alterations appeared as clustered mutations interspersed with polypeptide segments that were highly conserved. The regions of highest variability (dark shading in Figure 2) and highest conservation (light shading in Figure 2) were identified both by visual inspection and by objective determination of relative variation using computer analysis (Figure 3).

Predicted Secondary Structure

Since the extracellular region of the HTLV-III/LAV envelope glycoprotein constitutes a major target of the host immunologic response (Barin et al., 1985a), we examined this region for predicted antigenic epitopes using a computer program that predicts the secondary structure of proteins superimposed with values for hydrophilicity (Chou and Fasman, 1974; Hopp and Woods, 1981). Such an analysis of other proteins, including viral envelopes, has shown that antigenic epitopes are often associated with hydrophilic protein domains containing β turns (Atassi, 1978; Cohen et al., 1984; Westhoff et al., 1984; Eisenberg et al., 1985; Gunn et al., 1985; Pellett et al., 1985). This analysis demonstrated that the exterior envelope proteins of the five AIDS retroviruses each contain a number of sites that meet criteria for likely antigenic epitopes and that these regions generally coincide with the variable regions identified independently by amino acid sequence comparisons. Figure 4 illustrates the substantial differences in predicted secondary protein structure for the first variable region of the external envelope gene product for the five viruses. In this region, there were 7 β turns in HAT-3, 0 β turns in WMJ-1, 0 β turns in BH-10, 11 β turns in LAV-1a, and 5 β turns in ARV-2. The five envelope sequences also differed in this region in their degrees of hydrophilicity and in the number and location of potential N-linked glycosylation sites (Figures 2 and 4). In each of the other variable regions substantial differences existed similarly in predicted hydrophilicity, second-
Figure 1. Nucleotide Sequences of the Entire Envelope Genes of Five Independent AIDS Virus Isolates

Sequence information for BH-10 (HTLV-III), ARV-2, and LAV-1a was obtained from GENBANK. Sequences for WUJ-1 and HAT-3 were determined as part of the present study. Alignment of the sequences was performed pairwise with the assistance of NUCALN (Wilbur and Lipman, 1985). Numbering of nucleotides is from the first nucleotide of BH-10 following the TAG stop codon, and genomic regions corresponding to the signal peptide, extracellular glycoprotein (gp120), and transmembrane glycoprotein (gp41) are shown. Dashes indicate nucleotide identity with BH-10 and spaces indicate the absence of that nucleotide. The boxed nucleotides comprise adjacent perfect or imperfect direct repeats.
Figure 2. Amino Acid Sequences of the Entire Envelope Genes of Five Independent AIDS Virus Isolates

Sequence information for BH-10 (HTLV-IIib), ARV-2, and LAV-1a was obtained from GENBANK and information for WMJ-1 and HAT-3 from the present study. Alignment of the sequences was performed pairwise with the assistance of PRTALN (Wilbur and Lipman, 1983). Numbering of amino acids is from the first amino acid of BH-10 and regions corresponding to the signal peptide, extracellular envelope glycoprotein (gp120), and transmembrane glycoprotein (gp41) are shown. Dashes indicate amino acid identity with BH-10 and spaces indicate the absence of that amino acid. Arrows denote cysteine residues and solid and open circles denote conserved and nonconserved sites of potential N-linked glycosylation, respectively. Darkly shaded regions within the extracellular envelope glycoprotein correspond to regions of hypervariability as determined by both visual inspection and by computer analysis of variation (see Figure 3). Lightly shaded regions correspond to areas that are relatively highly conserved. Note that additional regions of intermediate variability are interspersed among highly variable and highly conserved areas.

ary structure, and potential glycosylation sites (S. Modrow, unpublished data).

Interspersed with the variable regions of the exterior envelope protein were other areas that were highly conserved among all five isolates analyzed. Most of these regions were primarily hydrophobic and contained only few or no \( \beta \) turns. Accordingly, they are believed to be less likely to represent antigenic epitopes. There were, however, a number of exceptions. One such exception is a conserved stretch of 46 amino acid residues immediately adjacent to the processing site of the envelope precursor (amino acids 473 through 518 in Figure 2). This area, shown also in Figure 4, was very highly conserved among all five viruses, contained numerous \( \beta \) turns, and was hydrophilic. Thus this conserved region of the exterior envelope glycoprotein would be expected to be both antigenic and cross-reactive among different viral strains. It has recently been shown that a synthetic peptide generated from a portion of this region does in fact detect naturally occurring antibodies in many patients infected with HTLV-III/LAV and that monoclonal antibodies directed against peptides of this region recognize the native gp120 (T. Palker and B. Haynes, personal communication).

Discussion

A growing body of scientific evidence indicates that the rate of evolution for RNA viruses in general is much
greater than for most prokaryotic or eukaryotic DNA genomes (for review, see Holland et al., 1982). From the data reported here, it is apparent that substantial heterogeneity exists among independent isolates of the AIDS virus. In another study (Hahn et al., 1986), we examined serial isolates of HTLV-III/LAV from chronically infected individuals and showed that the types of nucleotide changes described in this paper for independent virus isolates are also present in serial virus isolates from the same patients and that these changes occur over a very short period of time during the course of viral infection. In fact, we estimate the rate of genetic change for the HTLV-III/LAV envelope gene to be at least $10^{-3}$ nucleotide substitutions per site per year, a mutational rate equal to that of influenza A virus and a million times greater than for most eukaryotic genes (Hahn et al., 1986).

An understanding of the mechanisms by which genetic variation occurs in HTLV-III/LAV can be approached by comparative analysis of the five sequenced genomes described herein. The finding of frequent point mutations throughout the viral genome, and the analogy with other RNA viral systems (Zarling and Temin, 1975; Clements et al., 1980; O’Rear and Temin, 1982; Shivelman et al., 1983; Darlix and Spahr, 1983; Bruck et al., 1984; Salinovich et al., 1986), suggest that mutational events resulting from RNA-dependent DNA polymerase (reverse transcriptase) misreading coupled with a lack of proofreading enzymes contributes in a major way to genetic variation in HTLV-III/LAV. In addition to this, the presence of numerous adjacent perfect and imperfect direct repeats within the hypervariable regions of the external envelope gene (see boxed nucleotides in Figure 1) suggests that duplications, insertions, and deletions also contribute to genetic variation in HTLV-III/LAV. In this light, it is noteworthy that in some instances HTLV-III/LAV sequences appear to have been duplicated and then one of the copies changed, the result being that novel amino acids are introduced within that region. Direct repeats encompassing deletions have been observed in other biologic systems, both eukaryotic (Efstatiadis et al., 1980) and prokaryotic (Farabaugh et al.,...
1978), and in the lac I gene of E. coli such direct repeats correspond to hotspots of spontaneous genetic mutation (Farabagb et al., 1978).

Other potential mechanisms for genomic change in HTLV-III/LAV include recombination between different viral DNA molecules or between viral DNA and host DNA. Both processes are known to occur in other retroviral systems (Coffin, 1979; O’Rear and Temin, 1982; Donner et al., 1985; Bishop and Varms, 1985). In this study, and in a related study in which serial AIDS virus isolates from three patients were analyzed by restriction mapping or DNA sequencing (Hahn et al., 1986), evidence for recombination between different viral molecules was not apparent. However, given the high rate of genetic change in HTLV-III/LAV and the persistence within a given individual of more than one predominant viral form (Shaw et al., 1984; Wong-Staal et al., 1985; Hahn et al., 1986), genetic recombination between different HTLV-III/LAV molecules is a distinct possibility. The second process, recombination between viral and cellular DNA, appears much less likely. None of the AIDS virus isolates studied here contained DNA sequences closely related to normal human DNA as determined by Southern blot hybridization (Hahn et al., 1984; Luciw et al., 1984; Alizon et al., 1984), and comparison of their envelope nucleotide sequences did not reveal any large unexplained insertions or rearrangements (see Figure 1). Still another potential source for genetic variation in HTLV-III/LAV is the formation of stable secondary structures (stem–loops) in the viral RNA which in turn could lead to localized areas of reverse transcriptase misreading. This process has been proposed to explain in part the variation observed in foot and mouth disease virus (Weddell et al., 1985), but for HTLV-III/LAV we found no evidence for such stem-loop structures. In sum, the nucleotide sequence data from independent HTLV-III/LAV isolates, along with the analysis of sequential viral isolates from individual patients (Hahn et al., 1986), suggest strongly that point mutations, in combination with short deletions or insertions/duplications perhaps related to copy-choice misreading by the viral polymerase, are the primary mechanisms by which variation is generated in the AIDS virus.

The finding of genomic variation primarily in the extracellular envelope gene in regions having properties predictive of antigenicity raises the possibility that host-related immunologic pressures may select for variant viral strains. For two retroviruses related to HTLV-III/LAV, namely equine infectious anemia virus (EIAV) and visna virus, there is evidence that progressive changes in the envelope genes do in fact lead to substantial changes in envelope antigenicity (Clements et al., 1980; Montelaro et al., 1984; Salinovich et al., 1986). These genetic changes are localized predominately within the envelope gene, in contrast to gag and pol, and there is evidence that they result from immunologic selective pressures exerted by the host. For EIAV, there is considerable support for the idea that such changes in viral antigenicity are directly responsible for the chronic, periodic nature of the disease (Salinovich et al., 1986). For visna virus, similar changes in envelope antigenicity leading to loss of virus neutralization by host antibodies have been noted (Clements et al., 1980). However, the biologic relevance of these findings has been questioned by other workers who have discovered spread of virus in blood and CSF unabated by neutralizing antibodies (Petursson et al., 1976), long-term persistence of parental viral strains even after the appearance of variants (Lutley et al., 1983), and lack of antigenic variants in some animals even during advanced stages of disease (Thormar et al., 1983). Given the complexities of chronic retroviral-induced disease (Haase et al., 1977; Brahic et al., 1981; Stowring et al., 1985), it is likely that intrinsic viral characteristics as well as host-viral immune interactions play important roles in clinical outcome.

For HTLV-III/LAV, the biologic significance of envelope variation has been particularly difficult to ascertain because the nature of the host immunologic response to the virus has yet to be well characterized. Naturally occurring, cross-reactive, neutralizing antibodies specific for HTLV-III/LAV have been identified in individuals infected with HTLV-III/LAV (Robert-Guroff et al., 1985; Weiss et al., 1985). Furthermore, preliminary data also indicate that type-specific anti-HTLV-III/LAV antibodies are generated in vivo and that variants of HTLV-III/LAV evolve which, determined by their susceptibility to neutralization by homologous sera are antigenically distinct (M. Robert-Guroff and W. Parks, personal communication). Whether or not such type-specific neutralizing antibodies select for biologically important antigenic variants of HTLV-III/LAV during persistent viral infection in man remains to be seen.

Other mechanisms, unrelated to the immune system, could conceivably be involved in the selection of HTLV-III/LAV variants. For example, host-specific and tissue-specific properties of the cell membrane receptor for the virus could select for alterations in the viral envelope as the virus spreads through genetically different individuals or through different tissues within a given host (Shaw et al., 1985; Ho et al., 1985). Such a mechanism has been proposed as a possible explanation for envelope divergence in foot and mouth disease virus (FMDV), since this virus naturally infects many different animal species in-
cluding cattle, swine, sheep, goats, buffalo, and antelope (Wedell et al., 1985). A selection process for envelope variation related to cell membrane characteristics within and between species is also of special relevance to HTLV-III/LAV research, as are mechanisms involving genetic recombination, since a novel simian retrovirus (STLV-III, for Simian T-Lymphotropic Virus Type III) recently isolated from four diseased rhesus macaques (three with immunodeficiency and one with transmitted lymphoma) has been found to be very similar to HTLV-III/LAV in its morphology, protein structure, and immunologic characteristics (Kanki et al., 1985; Daniel et al., 1985). Antibodies to this virus (or a close relative) were identified in 28 of 67 healthy African green monkeys from central Africa (caught in the wild), as well as in a number of clinically asymptomatic West African prostitutes (Barin et al., 1985). The fact that this retrovirus has infected two species of Old World primates, and possibly man, and that it is highly related to HTLV-III/LAV suggests that the evolutionary origins of STLV-III and HTLV-III/LAV are similar. That their envelope proteins are less related than their gag proteins underscores the potential importance of variation in the envelope gene of HTLV-III/LAV as a determinant of biologic activity.

Whether or not the observed propensity for variation in HTLV-III/LAV has biologic significance has yet to be formally proven. However, such extensive variation in the different genes of this virus clearly has the potential for drastically altering both its immunologic and biologic properties. These include the antigenicity and tissue tropism conferred by env; the polymerase, protease, and integrase functions of pol; the gene regulatory properties ofLTR andtat; and other potentially important products of sor andorf whose functions are still unknown. Future efforts to understand viral pathogenesis and to develop effective preventive and treatment measures for AIDS will have to take into account the extensive genomic heterogeneity present in HTLV-III/LAV.

Experimental Procedures

Virus isolation

Virus was isolated from patients' peripheral blood mononuclear cells, transmitted to immortalized T cell lines, and propagated in tissue culture as described (Popovic et al., 1984; Hahn et al., 1986).

Molecular Cloning of HAT-3 and WMJ-1

Lambda (λ) phage libraries were constructed according to standard procedures (Maniatis et al., 1982) using the cloning vector λgtWes λB and Sat digested viralcellular DNA which had been enriched by sucrose gradient centrifugation for 8-10 kb fragments. Both HTLV-III/LAV and HTLV-IIIMet DNA contained a single 9 kb SstI fragment detectable using these subgenomic clones. Detailed restriction cleavage patterns were performed with a second enzyme, separated by polyacrylamide gel electrophoresis, and eluted into buffer. In some instances, double-labeled fragments were prepared, strand-separated, and sequenced individually. The sequences of both envelope genes and most of the remainder of HTAT-3 was confirmed by sequence analysis of both DNA strands. The computer program of Queen and Korn (1983) was used for transcribing the nucleotide sequences into amino acids. Nucleotide and amino acid sequences were aligned pairwise using NUCALN and PRTALN (Wilbur and Lipman, 1983).

Nucleotide Sequence Analysis

Nucleotide sequencing was performed according to Maxam and Gilbert (1980). Plasmid subclones were cleaved with the appropriate restriction enzymes and end-labeled using polynucleotide kinase and 32P-ATP for 5' ends, and a 32P-dNTP plus DNA polymerase (Klenow fragment) or a 39-end ATP plus terminal deoxynucleotidyl transferase for 3' ends. Labeled fragments were eluted with a cacodylate buffer and electrophoresed on a polyacrylamide gel. Sequence reactions were then performed using these subgenomic clones. Details of restriction cleavage patterns of both HAT-3 and WMJ-1 corresponded exactly (25 out of 25 restriction sites) to the predominant proviral forms present in the respective cell lines.

Acknowledgments

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