Human endogenous retroviruses (HERVs) are a family of viruses within our genome with similarities to present day exogenous retroviruses. HERVs have been inherited by successive generations and it is possible that some have conferred biological benefits. However, several HERVs have been implicated in certain cancers and autoimmune diseases. This article demystifies these retroviruses by providing an insight into HERVs, their means of classification, and a synopsis of HERVs implicated in cancer and autoimmunity. Furthermore, the biological roles of HERVs are explored.

H
m
d
Human endogenous retroviruses (HERVs) represent footprints of previous retroviral infection and have been termed “fossil viruses”. They are transmitted vertically through the germline and are thus inherited by successive generations in a Mendelian manner. Over time, HERVs have been subjected to repeated amplification and transposition events giving rise to multi-copy and single copy proviruses that are distributed within the DNA of all cells. Overall, HERVs constitute about 1% of the human genome. HERVs possess a similar genomic organisation to present day exogenous retroviruses such as human immunodeficiency virus (HIV) and human T cell leukaemia virus (HTLV), and are composed of gag, pol, and env regions sandwiched between two long terminal repeats (LTRs) (fig 1). The LTRs possess nucleotide sequence motifs that are fundamental to the regulation of retroviral gene expression. In brief, the gag and env genes encode retroviral capsid and envelope proteins, respectively, whereas the pol gene encodes enzymes for viral replication, integration, and protein cleavage. Retroviruses in effect are retrograde, because the flow of genetic information is reversed compared with the normal pathway of molecular biosynthesis—DNA → RNA → protein. Indeed, all retroviruses necessitate the conversion of viral RNA into cDNA intermediaries, which is catalysed by the enzyme reverse transcriptase. “Overall, human endogenous retroviruses constitute about 1% of the human genome”

Over 20 HERV families have been identified during the past two decades. Over 20 HERV families have been identified during the past two decades. Although many are defective through the accumulation of mutations, deletions, and termination signals within coding sequences, a limited number of HERVs have the potential to produce viral products and, indeed, to produce viral-like particles. Furthermore, some HERVs have been implicated in certain autoimmune diseases and cancers and might have a role in the aetiology and pathology of disease. However, many HERVs have been present in our genome for a considerable period of time so that their presence may also be of benefit to the human host.

MOLECULAR BIOLOGY AND HERVS
The focus of the human genome project is to sequence the 3 billion DNA bases that compose our human genome. Interestingly, only a small proportion (about 3%) appears to constitute the estimated 24 179 to 87 720 genes that are translated into the proteins necessary to life. The function of non-coding DNA (malign as “junk DNA”) is not readily obvious and it may have important roles in packaging and influencing gene expression. DNA is composed of two helical strands orientated in an antiparallel fashion (fig 2), where each strand consists of a deoxyribose phosphate backbone and a series of purine and pyrimidine bases denoted A (adenine), G (guanine), T (thymine), and C (cytosine). A unit of sugar, phosphate, and base is strictly termed a nucleotide, although casually represented by its base. The two DNA strands associate through the complementary pairing of bases—for example, A–T and C–G—which is stabilised through hydrogen bonding. Stretches of DNA, measured in base pairs, constitute the numerous genes that are found on the 23 pairs of human chromosomes. Genes vary in size, with some in the order of 20–40 kilobase pairs (kbp), whereas others extend to millions of base pairs. In addition, human genes are composed of exons, which are transcribed and translated into amino acids, and introns, which are interspersed between exons and represent non-translated regions that contribute to the large size of some genes. Retroviruses are remarkably small (in the order of 9–10 kbp) compared with other pathogenic viruses, such as cytomegalovirus, which is 200 kbp.

See end of article for authors’ affiliations

Correspondence to:
Dr P N Nelson, Molecular Immunology Laboratories, School of Applied Sciences, University of Wolverhampton, Wulftuna Street, Wolverhampton WV1 1SB, UK; P.N.NELSON@wlv.ac.uk

Accepted for publication
19 September 2002

Abbreviations:
CFTR, cystic fibrosis gene; CTL, cytotoxic T lymphocyte; EBV, Epstein-Barr virus; HERV, human endogenous retrovirus; HIV, human immunodeficiency virus; HTLV, human T cell leukaemia virus; IDDM, insulin dependent diabetes mellitus; IS, immunosuppressive peptide; LINES, long stretches of related sequences; LTR, long terminal repeat; MHC, major histocompatibility complex; MMTV, mouse mammary tumour virus; PCR, polymerase chain reaction; PTN, pleiotrophin gene; RA, rheumatoid arthritis; SLE, systemic lupus erythematosus; SS, Sjogren’s syndrome; TGCT, testicular germ cell tumour

www.molpath.com

Downloaded from http://mp.bmj.com/ on October 10, 2017 - Published by group.bmj.com
Nevertheless, retroviruses capitalise on their limited size by using frame shifts to retrieve genetic information from multiple overlapping open reading frames (fig 3). Interestingly, the open reading frames of many defunct HERVs are interrupted by premature stop codons that interrupt the “read through” of genetic information. A previous demystified article highlights the use of the polymerase chain reaction (PCR) to detect certain nucleotide regions of HERVs that are common to all retroviruses.

“Retroviruses capitalise on their limited size by using frame shifts to retrieve genetic information from multiple overlapping open reading frames”

Before protein synthesis, mRNA is transcribed from relevant stretches of DNA (reliant on a triplet base code or codon) that specifies a start signal (for example, ATG), an amino acid, or a termination signal (TAA, TAG, or TGA). The primary mRNA transcript is refined by splicing out unwanted introns and then by adding a series of adenine bases to its 3’ end. The process of polyadenylation is considered crucial to the stability of the mRNA that leaves the nucleus to attach to cytoplasmic ribosomes. Here, individual tRNA molecules attached to a specific amino acid are aligned with mRNA that dictates the assembly of polypeptide chains (fig 4). Strictly speaking, tRNAs use anticodons for complimentary binding, although a codon, such as AAU for lysine (denoted by K in the single amino acid code), is often used with reference to the tRNA “primer binding site” of a HERV—for example, HERV-K. Interestingly, tRNAs may use the “wobble” phenomenon in that the first two bases are essential for binding but the third base is less crucial. Therefore, it is possible that tRNAs may over-ride stop codons and allow the growth of a polypeptide chain. Ultimately, both exogenous and endogenous retroviruses exploit the cell’s genetic machinery and comply with the same generic principles of protein biosynthesis to produce further viral progeny. However HERVs stop short of viral budding (fig 3) and consequently are non-infectious.

CLASSIFICATION OF HERVS

Historically, retroviruses were shown to cause malignant tumours and leukaemias in chickens and mammals—for example, the Rous sarcoma virus and mouse mammary tumour virus (MMTV), respectively—and were subdivided into three subfamilies: oncovirinae (oncoviruses), lentivirinae (slow viruses), and spumavirinae (foamy viruses). Retroviruses were also classified according to morphological and biological criteria: in particular by the observance of retroviral particles in infected cells. In brief, type A retroviruses were only visible inside cells, devoid of envelope, and referred to as intracisternal particles (A type particles), whereas types B, C, and D were enveloped and produced extracellular particles that varied in size and appearance. Subsequently, in the 1980s, the retroviruses HTLV-I and HIV-1 were reported and a new classification formulated by the International Committee for Taxonomy of Viruses. This taxonomy reflected the similarity of a virus to an established retroviral genus, such as murine leukaemia related virus. Set against this background, the first HERV was reported in 1981, and many others have subsequently been identified (table 1) using several different methodologies. These have included the screening of human

**Figure 1** Salient features of human endogenous retroviruses (HERVs) and similarity to exogenous retroviruses.

| HERVs: footprints of previous exposure to retroviruses; coined “fossil viruses” | Constitute approximately 1% of the human genome |
| Similar genomic organisation to exogenous retroviruses e.g. HIV-1, HTLV-I |
| Transmitted vertically in the germline through successive generations |
| Possess a reverse transcriptase gene region found in all retroviruses |

**Figure 2** A schematic diagram of a segment of DNA that is transcribed into mRNA and finally translated into a peptide product. In this example the DNA triplet code AAG is translated into the amino acid lysine.

12 Nelson, Carnegie, Martin, et al
and env regions. Members include HERV-H, HERV-I, and HERV-R (ERV-9). Class II HERVs show homology to mammalian type B (for example, MMTV) and type D retroviruses and are subdivided into 10 groups. These groups reflect the sequence homology of reverse transcriptase regions of PCR derived clones from normal leucocytes, and clones derived by low stringency hybridisation of an MMTV gag–pol probe to a breast cancer cell genomic library. Members include HERV-K (fig 5) and HERV-K (C4). Interestingly, all class II HERVs possess a lysine tRNA reflecting their derivation from B and D type viruses. HERV-K has been termed the biologically most active human endogenous retrovirus family, and has been further subdivided into type 1 or type 2, based on the presence or absence of a 292 bp segment at the pol–env boundary. Finally, foamy virus related HERVs are classified as class III HERVs and include a solitary member, HERV-L. This system provides a uniformity to the classification of HERVs but does not include the human T cell leukaemia related endogenous retrovirus HRES-1, which shows only limited homology to HTLV-I in the LTR region.

**HERVS AND CANCER**

Although the precise role(s) of HERVs in the carcinogenic process has not been fully elucidated there are several studies that, if taken together, put forward a convincing argument for the possible involvement of HERVs in malignancy. HERVs may be involved in carcinogenesis by virtue of the expression of HERV mRNA, functional proteins, or retroviral-like particles. They may also be associated with the generation of new promoters or the activation of proto-oncogenes. The expression of HERV mRNA is increased in some cases of small cell lung carcinoma. In addition, a teratocarcinoma cell line has been shown to possess a HERV-K sequence and to secrete retroviral-like particles. Testicular germ cell tumours (TGCTs) have been shown to contain proteins of the HERV-K family and patients with TGCT often exhibit a specific immune response to gag and env proteins. It has been suggested that HERV-K may be important in the progression of TGCT through inhibition of an effective immune response, and the HERV env genes have been shown to encode immunosuppressive proteins. It is clear that overexpressed HERV proteins can elicit high titre IgG responses in some settings (for example, HERV-K10 in patients with renal cancer).
as detected by the SEREX method (serological identification of expressed genes),
suggesting that HERV proteins may in the future provide targets for antitumour
immunotherapy.

It has been suggested that HERV-K may be important in the progression of testicular germ cell tumours through inhibition of an effective immune response.

HERV-K might be important in the pathogenesis of human breast cancer. It has been shown that the T47D human mammary carcinoma cell line produces retroviral particles with reverse transcriptase activity. Both the HERV-K10 related sequences of T47D cells and the reverse transcriptase activity are increased by steroid hormone treatment, which is thought to be the result of transcriptional activation via binding of the progesterone receptor to regions on the HERV-K genome that correspond to progesterone and glucocorticoid response elements.

In choriocarcinoma, it has been shown that a HERV type C is inserted into the human growth factor gene, pleiotrophin (PTN). This results in the generation of a novel tissue specific promoter, which results in the expression of HERV–PTN fusion transcripts, leading to the production of biologically active PTN protein. Expression of the PTN protein (which is normally expressed only at very low amounts in a few normal adult tissues) appears to be responsible for the aggressive and invasive growth of human choriocarcinoma.

Overexpression is a common mechanism by which proto-oncogenes become activated, leading to subsequent neoplastic transformation. In particular, activation of proto-oncogenes of the ras family is common in many tumour types, and some studies have suggested a potential role for HERV's in ras activation. It was shown a methylaminoacetate induced rat mammary carcinoma that insertion of a defective endogenous retrovirus into the intron of c-Ha-ras was responsible for its more than 10 fold overexpression.

Recently, it has been shown that lymphotrophic herpesvirus, Epstein-Barr virus (EBV), itself a potent transforming agent, can transcriptionally activate the env gene of HERV-K18, which possesses superantigen activity (as demonstrated by major histocompatibility complex (MHC) class II dependent preferential activation of TCRVB13 T cells in response to murine B cells transfected with the HERV-K18 env gene). The authors suggested that this phenomenon accounts for the previously described EBV associated superantigen activity, which might in turn be crucial to T cell activation by EBV. Although the exact function of HERV's in the carcinogenic process is still under investigation, the evidence implicating HERV's in the carcinogenic process is substantial and further investigation will be required to elucidate the contribution of HERV's to the development of malignancy.

HERVS AND AUTOIMMUNITY

In 1990, an article appeared in the Times newspaper (24 November) with the title "AIDS-like virus may cause arthritis". The report focused on Robert Garry's research that identified retroviral particles in lip biopsies taken from patients with primary Sjogren's syndrome (SS). Similarly, in other autoimmune rheumatic diseases, such as rheumatoid arthritis (RA) and systemic lupus erythematosus (SLE), a plethora of articles added to this intriguing observation by providing evidence of retroviral antigens at the site of disease, or the presence of antiretroviral antibodies in the sera of patients.

A novel report in 1994 used both PCR (using consensus primers) and serological tests to investigate the presence of retroviruses in a cross section of patients with rheumatoid diseases, including RA, SS, and SLE. Interestingly, PCR failed to amplify products relating to HTLV-I or HIV-1, although antibodies to retroviral antigens were present but no evidence to implicate exogenous retroviruses. However, in the case of rheumatoid arthritis (RA), PCR could detect proviral DNA of the HML-1 family (HERV-K) in 60% of the patients, whereas only 18% of healthy controls were positive.

A novel report in 1994 used both PCR (using consensus primers) and serological tests to investigate the presence of retroviruses in a cross section of patients with rheumatoid diseases, including RA, SS, and SLE.

Table 1. Classification of human endogenous retroviruses (HERVs)

<table>
<thead>
<tr>
<th>HERV family</th>
<th>Representative accession number</th>
</tr>
</thead>
<tbody>
<tr>
<td>Class I HERVs (type C related HERVs)</td>
<td></td>
</tr>
<tr>
<td>Group 1, HERV-HF</td>
<td></td>
</tr>
<tr>
<td>HERV-H (RTLVh, RGH)</td>
<td>AF108842</td>
</tr>
<tr>
<td>HERV-F</td>
<td>AF070684</td>
</tr>
<tr>
<td>Group 2, HERV-RW</td>
<td></td>
</tr>
<tr>
<td>HERV-W</td>
<td>AF072506</td>
</tr>
<tr>
<td>HERV [ERV]</td>
<td>X57147</td>
</tr>
<tr>
<td>HERVP (HuERS-P, HuRRS-P)</td>
<td>X06279</td>
</tr>
<tr>
<td>Group 3, HERV-E</td>
<td></td>
</tr>
<tr>
<td>HERVE [81, ERVɑ, NP2*]</td>
<td>S46403</td>
</tr>
<tr>
<td>51:1</td>
<td>J00273</td>
</tr>
<tr>
<td>HERV [ERV3]</td>
<td>M12140</td>
</tr>
<tr>
<td>HERV [ERV4]</td>
<td>M64936</td>
</tr>
<tr>
<td>Group 4, HERV-T</td>
<td>M32788</td>
</tr>
<tr>
<td>HERV T [S71, CRTK1, CRTK6]</td>
<td></td>
</tr>
<tr>
<td>Group 5, HERVP</td>
<td></td>
</tr>
<tr>
<td>HERV [RTLV4]</td>
<td>X14953</td>
</tr>
<tr>
<td>HERVP [ERV-FD]</td>
<td>U27241</td>
</tr>
<tr>
<td>Group 6, ERV-FRD</td>
<td></td>
</tr>
<tr>
<td>ERV-FRD</td>
<td>U27240</td>
</tr>
<tr>
<td>Class II HERVs (type A, B, and D related HERVs)</td>
<td></td>
</tr>
<tr>
<td>Group 1, HERV [HML-1]</td>
<td></td>
</tr>
<tr>
<td>HERV [HML-1]</td>
<td>U35102</td>
</tr>
<tr>
<td>HERV [HML-2]</td>
<td>M14123</td>
</tr>
<tr>
<td>HERV-HTDV</td>
<td>X8227</td>
</tr>
<tr>
<td>Group 3, HERV [HML-3]</td>
<td></td>
</tr>
<tr>
<td>HERV K [HML.3]</td>
<td>U35153</td>
</tr>
<tr>
<td>Group 4, HERV [HML-4]</td>
<td></td>
</tr>
<tr>
<td>HERV-K [HML-5]</td>
<td>AF020092</td>
</tr>
<tr>
<td>HERV K [HML-6]</td>
<td>AF015995</td>
</tr>
<tr>
<td>HERV KNMMW2</td>
<td></td>
</tr>
<tr>
<td>HERV K [HML-6p]</td>
<td>U86698</td>
</tr>
<tr>
<td>Group 6, HERV KNMMW2</td>
<td></td>
</tr>
<tr>
<td>HERV K [HML-7]</td>
<td>AF016000</td>
</tr>
<tr>
<td>HERV KNMMW7</td>
<td>AF015996</td>
</tr>
<tr>
<td>Group 7, HERV KNMMW3</td>
<td></td>
</tr>
<tr>
<td>HERV K [HML-9]</td>
<td>AF016001</td>
</tr>
<tr>
<td>HERV KNMMW9</td>
<td></td>
</tr>
<tr>
<td>Group 8, HERV [HML-8]</td>
<td></td>
</tr>
<tr>
<td>HERV KNMMW3</td>
<td>AF016000</td>
</tr>
<tr>
<td>Group 9, HERV [HML-9]</td>
<td></td>
</tr>
<tr>
<td>HERV KNMMW9</td>
<td>U07856</td>
</tr>
<tr>
<td>HERV-KC4</td>
<td></td>
</tr>
<tr>
<td>Group 10, HERV [HML-10]</td>
<td></td>
</tr>
<tr>
<td>HERV-KC4</td>
<td>X89211</td>
</tr>
</tbody>
</table>

Table 1 Classification of human endogenous retroviruses (HERVs)
of neo-antigens, perhaps as a result of HERV and/or exogenous viral combinations. The use of animal models has also served to enhance our understanding of endogenous retroviruses. In a lupus model, an 8.4 kbp endogenous retroviral transcript is expressed in affected mice. Furthermore, a retroviral element in one of the introns of the fas apoptosis gene appears to alter the splicing of fas transcripts, resulting in a lupus-like autoimmune disease in MRL-lp/lpr mice. Further investigations using animal models and multi-centre patient studies are needed to establish links between specific HERVs and autoimmune diseases because many HERVs are also expressed in varying amounts, or in a coordinated fashion, in normal tissues.

“Mechanisms whereby HERVs could influence autoimmunity include molecular mimicry, superantigen motifs that bypass the normal MHC restrictive process of T cell stimulation, aberrant expression of antigens, and the presence of neo-antigens”

**BIOLOGICAL IMPORTANCE OF HERVS**

Phylogenetic studies have shown that some HERVs emerged over 3 million years ago, whereas others appeared after the divergence of hominoid and ape lineages. Consequently, HERVs have been present in our genome for a considerable period of time and perhaps have been retained because they performed a useful biological function. Alternatively, some HERVs may have been difficult to eliminate and thus persisted during evolution. To confer a selective advantage, a premise remains that HERVs produce products and/or augment mechanisms that benefit host survival. Several possibilities are highlighted below.

**Immunosuppressive peptide**

The product derived from the env gene of mammalian type C retroviruses possesses motifs—for example, the fusion peptide, leucine zipper protein, and immunosuppressive peptide (ISP)—that are essential for fusion and the infection of cells. In brief, the precursor env product is cleaved into two components: a surface protein (gp70) and a transmembrane protein (p15E) that contains an immunosuppressive region. Of interest is an ISP, termed CK5-17, that suppresses lymphocytes and alters cytokine profiles in animal models. The ISP sequence from the murine leukaemia virus LQNRRGLDLLFLKEGGLC (single amino acid code) is reasonably well conserved in HERV-H19 (LQNRRGLDLTAEGGGLC), HERV-R (ERV-3) (YQNRLADYLLAEEGVC), and HERV-E(4–1) (YQNRLADYLLAAEGGVC), but less so in HERV-K10 (FEASKAHLNVPGEAIA). The presence of an ISP could be advantageous to a virally infected cell—in terms of shielding or “cloaking” itself from immunological attack—but may equally be important to a host. This is perhaps exemplified by HERV-R (ERV3), which is highly expressed in trophoblastic cells and results in high concentrations of env protein (~65 kDa) in syncytiotrophoblasts. The immunosuppressive potential of this HERV and the fusogenic nature of placenta tissue suggests a possible involvement in normal placental function, in protecting the developing fetus from maternal
immune responses. Furthermore, it is possible that HERVs may change the pattern of gene expression during embryo development by altering different rates of development of different parts of the embryo. Another endogenous retrovirus, HERV-W, has also been shown to encode a protein termed syncytin, which may have a role in placental morphogenesis.

**Antiviral resistance**

Mechanisms of conferring protection against a related viral agent may include retroviral receptor blockade (by HERV products) and interference of replication by antisense mRNA. However, it is also possible that HERV peptides could prime an immune response to an undesirable agent. Intriguingly, cohorts of female sex workers in the Gambia and Kenya have been identified who remain uninfected and seronegative despite being repetitively exposed to HIV. The apparent “resistance” to HIV infection is suggested to result from MHC class Irestricted cytotoxic T lymphocytes (CTLs)—HLA-B35 and HLA-A2 (A*6802)/HLA-B18—which are found in Gambians and Kenyans, respectively. The immune response to HIV infection is characterised by a vigorous HIV specific CTL response where virus-specific CTLs recognise antigen in the form of processed peptides (eight to 10 amino acids in length), which are bound in the cleft of MHC class I molecules on the surface of antigen presenting cells. In this context, recent data have highlighted similarities between HIV CTL peptide sequences and regions of HERV-K10. Consequently in some cases previous exposure to HERV peptides could potentially immunise certain individuals, although this argument does not explain why some females seroconvert after a reduction in sex work.

**Long terminal repeats**

Retroviral genes that have been integrated into the genome are bordered by short direct repeats of host DNA and LTR sequences of about 500–600 nucleotides. These LTRs can influence neighbouring genes because they may contain transcriptional regulatory elements such as enhancers, promoters, hormone responsive elements, and polyadenylation signals. The influence on protein expression is demonstrated by proteins such as salivary amylase, ZNF80, cytochrome C1, Kruppel-like H-plk, and phospholipase-A2, whose genes are linked to LTRs. Numerous solitary LTRs are also present within our genome. These sequences, which include about 10 000 copies of solitary LTRs pertaining to HERV-K, have arisen through recombination events that could have “deleted” harmful retroviral genes. It is noteworthy that LTRs are located near MHC genes and thus could modulate antigen expression, particularly in individuals who possess identical MHC genes but different LTRs. With reference to autoimmune diseases, the presence of an HERV-K LTR in the HLA-DQ region (DQ-LTR3) has been shown to have a strong influence on the occurrence of IDDM. Furthermore, the presence of DQ-LTR on HLA-DQB1*0302 and its absence on DQB1*0201 are independent risk markers for IDDM. The presence of DQ-LTR has also been shown to be a marker for increased susceptibility to RA in subjects with the HLA-DR4-DQB1*0301 and HLA-DR4-DQB1*0302 haplotypes. In addition, polymorphisms within LTRs could be important because changes in sequence could greatly affect promoter activity. Interestingly, three allelic forms of the LTR of HERV-R (ERV3) have been shown, although no link could be associated with disease, in this case multiple sclerosis. The fact that LTRs interact with chemicals (such as carbon tetrachloride), hormones, and environmental agents and function as switches for alternative splicing (for example, the short and long forms of the human leptin receptor) highlights the importance of these elements. In addition, their location (adjacent to MHC or T cell receptor genes) and transactivation by helper viruses’ warrants further research into their potential role in autoimmune diseases and cancer.

**Plasticity**

HERVs constitute only a part of what are termed “transposable elements”, a generic term encompassing both DNA sequences that can be excised and reinserted at another site and retroelements. The term retroelements describes any sequence that can replicate itself by a process involving reverse transcription, and includes HERVs, retrotransposons (which mostly lack an env gene), retroprosomes, and retrosequences. Retroprosomes and retrosequences are exemplified by long stretches of related sequences (up to 6 kbp) called LINES and very short interspersed repeat elements of about 300 bp, respectively. Thus far, from being a fixed, immutable structure, the genome of a eukaryotic cell can harbour many sequences that move from one site on a chromosome to a completely different position. This phenomenon of plasticity is considered important because it permits rapid changes in our genome that could not be afforded by mutations alone. Furthermore, retroelements may carry regulatory sequences (such as enhancers and promoters) to new sites in the genome and thus alter the expression of existing adjacent genes. Of course, there remains a potential to disrupt genes by insertion mutation and produce defective and/or truncated products, as has been highlighted by Kazazian, who reported cases of human genetic disease caused by the random insertion of LINES.

**PREMATURE STOP CODONS**

It is known that many HERVs possess premature stop codons and of course the RNA products of truncated genes may well be eliminated before protein translation. However, examples of premature stop codons that result in human disease by the production of a truncated protein product have been demonstrated in cases of the DNA repair disorder xeroderma pigmentosum and also in leukaemia. In addition, approximately 5% of cystic fibrosis cases are caused by premature termination codons. Interestingly, some aminoglycoside antibiotics appear to suppress stop codons in several organisms. Howard and colleagues demonstrated this effect in cell cultures that expressed constructs carrying two different premature stop codons within the cystic fibrosis gene (CFTR). Treatment with antibiotic produced a full length CFTR: the physiological suppression of the premature stop codon was thought to result from a mispairing of an amino-acyl tRNA that successfully bound to a stop codon. The importance of this observation is that HERVs, despite possessing premature stop codons, may have the potential to produce truncated or full length products that are fundamental to the mechanisms of molecular mimicry, aberrant expression of products, and the stimulation of T cells through superantigen motifs.

**CONCLUSIONS**

This article has provided an overview of a complex topic that may have ramifications in host protection, cancer, and autoimmunity. Ultimately, are HERVs friends or foes? In conferring a biological advantage, HERVs (and solitary LTRs) may indeed be beneficial. Their role in immunological homeostasis and perhaps protection against exogenous retroviruses is intriguing. Alternatively, HERV insertion mutation, molecular mimicry, superantigen motifs, and recombination with other viruses could be responsible for the development and pathology of disease. An additional aspect is whether the presence of HERV peptides during ontogeny culminates with a hole in the immune repertoire. As a result, peptides with similarity to HIV-CTL sequences could be more dangerous to a given individual.
REFERENCES


www.molpath.com


Demystified . . . Human endogenous retroviruses


Mol Path 2003 56: 11-18
doi: 10.1136/mp.56.1.11

Updated information and services can be found at:
http://mp.bmj.com/content/56/1/11

These include:

References
This article cites 80 articles, 23 of which you can access for free at:
http://mp.bmj.com/content/56/1/11#BIBL

Email alerting service
Receive free email alerts when new articles cite this article. Sign up in the box at the top right corner of the online article.

Notes

To request permissions go to:
http://group.bmj.com/group/rights-licensing/permissions

To order reprints go to:
http://journals.bmj.com/cgi/reprintform

To subscribe to BMJ go to:
http://group.bmj.com/subscribe/