Genome analysis

Retroviral promoters in the human genome

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ABSTRACT

Motivation: Endogenous retrovirus (ERV) elements have been shown to contribute promoter sequences that can initiate transcription of adjacent human genes. However, the extent to which retroviral sequences initiate transcription within the human genome is currently unknown. We analyzed genome sequence and high-throughput expression data to systematically evaluate the presence of retroviral promoters in the human genome.

Results: We report the existence of 51 197 ERV-derived promoter sequences that initiate transcription within the human genome, including 1743 cases where transcription is initiated from ERV sequences that are located in gene proximal promoter or 5′ untranslated regions (UTRs). A total of 114 of the ERV-derived transcription start sites can be demonstrated to drive transcription of 97 human genes, producing chimeric transcripts that are initiated within ERV long terminal repeat (LTR) sequences and read-through into known gene sequences. ERV promoters drive tissue-specific and lineage-specific patterns of gene expression and contribute to expression divergence between paralogs. These data illustrate the potential of retroviral sequences to regulate human transcription on a large scale consistent with a substantial effect of ERVs on the function and evolution of the human genome.

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Supplementary information: Supplementary data are available at Bioinformatics online.

1 INTRODUCTION

Approximately 5% of the human genome sequence is derived from retroviruses (Lander, 2001). Retroviral genomic sequences are remnants of past infections that resulted in the integration of provirus genomes into the DNA of germline cells (Bock, 2000; Bromham, 2002). The abundance of these so-called endogenous retrovirus sequences (ERV's) testifies to the extent that human evolution has been shaped by successive waves of viral invasion (Sverdlov, 2000).

One way that ERVs have affected the function and evolution of the human genome is by donating regulatory sequences that control the expression of nearby genes. The gene regulatory effects of ERVs were first uncovered in a number of anecdotal studies on specific genes (reviewed in Bannert et al., 2004; Medstrand et al., 2005). For instance, the long terminal repeat (LTR) of a human ERV (HERV-E) was shown to serve as an enhancer element that confers parotid-specific expression on the amylase gene (Samuelson et al., 1990).

Later, more systematic computational analyses of the human genome sequence revealed that many human genes contained ERV-derived regulatory regions, suggesting an even greater contribution of retroviruses to human gene regulation (Jordan et al., 2003; van de Lagemaat et al., 2003). Continued efforts to characterize ERV-derived promoters have turned up several new cases in recent years (Dunn et al., 2003, 2006; Romanish et al., 2007). Nevertheless, the full extent of the contribution of ERV sequences to the initiation of transcription in the human genome has yet to be appreciated.

Initiation of transcription by ERV promoters often results in the production of alternative transcripts that are both tissue-specific and lineage-specific. For instance, testis-specific expression of the human gene encoding the neuronal apoptosis inhibitory protein (NAIP) is driven by an LTR promoter sequence, whereas a distinct LTR promoter in rodents confers constitutive expression of the orthologous gene (Romanish et al., 2007). An ERV LTR sequence also serves as an alternative promoter that drives expression of the beta1,3-galactosyltransferase five gene specifically in colorectal tissue (Dunn et al., 2003).

The lineage-specific regulatory effects of ERV promoters can be attributed to the fact that ERV sequences result from past germline infections, many of which occurred relatively recently along specific evolutionary lineages. In fact, most of the ERV sequences in the human genome are primate-specific (Sverdlov, 2000), while most human genes are far more ancient and share orthologs with distantly related species (Lander et al., 2001). This means that regulatory effects exerted by ERV promoters will often lead to expression differences between primate and non-primate orthologs or between deeper evolutionary lineages for more ancient ERVs. In other words, ERV promoters are likely to drive evolutionary changes in gene expression, long thought to be an important determinant of species divergence (King et al., 1975).

The application of novel high-throughput techniques for the analysis of gene expression has revolutionized the study of the human transcriptome and revealed far more regulatory complexity than previously imagined. Two techniques in particular, cap analysis of gene expression (CAGE) and paired-end ditag (PET) sequencing, enable the precise genome mapping of many thousands of promoter sequences that initiate transcription. CAGE is a technique that allows for the characterization of short sequence tags from the 5′-most ends of full-length cDNAs (Shiraki et al., 2003). Accordingly, mapping CAGE tags to the human genome unambiguously identifies transcription start sites (TSS) and their corresponding promoters. PET sequencing involves the determination of sequences for tags from both the 5′ and 3′ ends of full-length cDNAs (Ng et al., 2005). Thus, when PETs are mapped to the genome,

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A total of 49,814 mapped CAGE tag clusters, each corresponding to Atlas version 2 (GNF2) (Su et al., 2004). Transcriptional units (TUs) are defined as genomic regions spanning the 5' and 3' ends of full-length transcripts, they can be used to identify transcripts that are initiated within ERV sequences and read-through into human gene sequences.

2 METHODS

CAGE tags (n = 1,551,672) were downloaded from the Japanese National Institute of Genetics website (http://genomenetwork.nig.ac.jp/public/downloads/cage_Database_e.html) and mapped to the human genome as previously described (Shiraki et al., 2003). The human genome locations of PETs (n = 669,840) were taken from the UCSC Genome Browser (Karolchik et al., 2003) annotations (http://www.genome.ucsc.edu/gold-bin/lifetracks/LifetrackU185/hg18/utrData/Multiplatform). The PETs were generated from several cell lines: log phase of MCF7 cells (113,858), HCT116 cells treated with 5-fluorocil (124,770) and log phase of embryonic stem cell hES3 (426,301). Overlapping CAGE tags and overlapping PETs were clustered to identify individual TSS on the human genome. The UCSC Table Browser (Karolchik et al., 2004) and the program Galaxy (Gardene et al., 2005) were used to compare the locations of CAGE tags and PETs to the locations of human ERVs annotated with the RepeatMasker program (Smitt et al., 1996–2004). Only ERVs' sense strands, as opposed to more ancient mammalian apparent LTR-retrotransposons (MaLR), were analyzed here. The National Center for Biotechnology (NCBI) Refseq (Pruitt et al., 2007) gene model annotations were used to evaluate the production of chimeric transcripts that are initiated by ERVs and read-through into human genes.

3 RESULTS AND DISCUSSION

A total of 49,814 mapped CAGE tag clusters, each corresponding to an individual TSS, were found to map to the ERV LTR sequences (Table 1). The high number of ERV-derived TSS in the human genome identified with CAGE tag mapping underscores the potential of retroviral promoters to drive transcription. However, it is not possible to directly assess whether retroviral promoters identified using CAGE tag mapping actually drive the expression of known human genes. In fact, most of the ERV promoters identified with CAGE map to intergenic regions. This intergenic ERV promoter activity is likely to be a relic of the ERVs' ability to drive transcription of their own genome sequences from LTR promoters and may not necessarily be related to the transcription of human genes. Nevertheless, the presence of widespread ERV promoter activity in the human genome demonstrates that ERV sequences can maintain the ability to promote transcription for millions of years after their initial insertion into the genome.

In addition to the intergenic ERV promoters, there are 9292 CAGE identified ERV promoters that initiate transcription within 1 kb upstream or downstream of the previously characterized TSS of known human genes (Table 2). 1550 of these ERV CAGE tag clustersmap to 5' UTRs, consistent with transcription from previously characterized promoters, but the majority (7742) map just upstream of the 5' UTR, in the proximal promoter region, or downstream within genes' TUs. Therefore, these ERV-derived promoters are likely to be responsible for generating alternative transcripts of human genes.

PET sequence mapping data were also used to search for transcripts that are initiated from ERV promoters, and there are 1513 cases of PET identified ERV promoters in the human genome (Table 1). Because PET sequence tags include both the 5' and 3' ends of full-length transcripts, they can be used to identify transcripts that are initiated within ERV sequences and read-through into human gene regions. These cases correspond to chimeric transcripts, composed partially of both ERV and human gene sequences, and demonstrate ERV promoted expression of human genes. This approach identified 114 distinct retroviral TSS that promote transcription of human genes (Table 2 and 3).

Table 1. Numbers of ERV-derived TSS in the human genome

<table>
<thead>
<tr>
<th>Data source</th>
<th>Total TSS&lt;sup&gt;b&lt;/sup&gt;</th>
<th>Gene-associated TSS&lt;sup&gt;b&lt;/sup&gt;</th>
</tr>
</thead>
<tbody>
<tr>
<td>CAGE</td>
<td>49,814</td>
<td>9292</td>
</tr>
<tr>
<td>PET</td>
<td>1,513</td>
<td>114</td>
</tr>
</tbody>
</table>

<sup>b</sup>Total number of tag clusters representing individual ERV-derived TSS.

Table 2. Numbers of ERV-human gene associated or chimerics transcripts

<table>
<thead>
<tr>
<th>PET&lt;sup&gt;a&lt;/sup&gt;</th>
<th>CAGE&lt;sup&gt;a&lt;/sup&gt;</th>
</tr>
</thead>
<tbody>
<tr>
<td>PET 3' ends</td>
<td>PET 5' ends</td>
</tr>
<tr>
<td></td>
<td></td>
</tr>
<tr>
<td>Upstream</td>
<td>5' UTR</td>
</tr>
<tr>
<td>TU</td>
<td>5</td>
</tr>
<tr>
<td>3' UTR</td>
<td>12</td>
</tr>
<tr>
<td>Downstream</td>
<td>4</td>
</tr>
<tr>
<td>5' UTR</td>
<td>11</td>
</tr>
</tbody>
</table>

<sup>a</sup>Counts for ERV-derived CAGE sequence tag clusters that map within 1 kb upsteam of human Refseq gene 5' UTRs or within 5' UTRs, consistent with transcription from previously characterized promoters, but the majority (7742) map just upstream of the 5' UTR, in the proximal promoter region, or downstream within genes' TUs. Therefore, these ERV-derived promoters are likely to be responsible for generating alternative transcripts of human genes.

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Retroviral promoters

Fig. 1. MER4A alternative promoter of the GSTO1 gene. (A) The MER4A (red) ERV sequence is located in the proximal promoter region <500 bp from the GSTO1 5′ UTR. The locations of PET sequences (green) and spliced ESTs (black) are shown. (B) The MER4A (red) sequence region is enlarged and the individual PET sequences (green) and spliced ESTs (black) that support the existence of this promoter are shown. (C) Evolutionary conservation of MER4A versus GSTO1. MER4A is only found in chimp and rhesus and no other mammals (green bars), i.e. it is not conserved, whereas the adjacent GSTO1 exons are conserved across mammalian species (green and blue bars).

Supplementary Table 1). Twenty-one of these retroviral promoters have colocated ESTs, which independently support their ability to initiate transcription. These retroviral TSS correspond to 124 Refseq transcripts over 97 distinct gene loci. The positions of TSS for ERV-derived human gene promoters were analyzed to evaluate whether ERVs provide canonical promoters or promote alternative transcripts. While there are a number of ERV TSS that map to 5′ UTRs (Table 2 and Supplementary Figure 1), and are thus taken to promote transcription at (or near) previously characterized TSS, the majority of ERV promoters promote alternative transcription of human genes from upstream regions or from within the TU (Table 2). This further underscores the fact that ERVs promote alternative transcription of human genes.

The ability of ERVs to promote alternative transcripts of human genes is illustrated (Fig. 1) by the case of an alternative promoter of the glutathione-s-transferase omega 1 encoding gene (GSTO1 Refseq accession NM_004832) found on chromosome 10q25.1. The GSTO1 protein is a member of the theta class glutathione s-transferase-like family, and it has been shown to act as a stress response protein through cellular redox homeostasis (Kodym et al., 1999). GSTO1 nucleotide polymorphisms have been implicated in a number of cerebrovascular diseases including Alzheimer’s disease, Parkinson’s disease, vascular dementia and stroke (Kolsch et al., 2004; Li et al., 2003).

There is an ERV LTR sequence from the MER4A subfamily of sequences <500 bp upstream of the Refseq annotated 5′ UTR of GSTO1 (Fig. 1A). There are 15 individual PET sequences, forming three distinct TSS clusters, that have 5′ ends inside of the MER4A sequence and 3′ ends in the 3′ UTR of GSTO1 (Fig. 1A and B). All of the MER4A PET sequences were derived from only one of the four PET libraries ($\chi^2 = 8.6, P = 0.04$), log phase of embryonic stem cell hES3, indicating that this promoter is tissue- or condition-specific. In addition to the PET sequence-based evidence, there are a number of spliced expressed sequence tags (ESTs) that also indicate the MER4A sequence as an alternative promoter for GSTO1 (Fig. 1B).

Inspection of multiple sequence alignments of complete mammalian genomes reveals that the GSTO1 adjacent MER4A insertion is present in the human, chimp (Pan troglodytes) and rhesus (Macaca mulatta) genome sequences but absent in the bushbaby (Otolemur garnetti), mouse (Mus musculus), rat (Rattus norvegicus) and all other placental mammal sequences (Fig. 1C). In other words, that particular MER4A sequence inserted after the primate radiation began, and it is specific to the Haplorrhini suborder, which includes both new world and old world monkeys. On the other hand, the adjacent exonic sequences of GSTO1 show marked conservation compared to MER4A (Fig. 1C). Comparative sequence analysis with BLASTP indicates that GSTO1 is far more ancient than the MER4A insertion, having well-conserved orthologs among mammals and other vertebrates along with Drosophila melanogaster, Caenorhabditis elegans and a number of other more distantly related species.

The comparative sequence analysis suggests the possibility that the MER4A insertion may confer lineage-specific expression pattern on GSTO1. Furthermore, there are two human paralogs of GSTO1, GSTO2 and the pseudogene GSTO3P1, neither of which has the upstream MER4A insertion. So the specific regulatory effects of
expression levels across tissues as does the human
not significantly different from 0 (P = 0.77) and correlation of
expression levels across tissues as does the human GSTO1
GSTO1 paralogs using microarray data from the Novartis Gene
Expression Atlas version 2 (GNF2) (Su et al., 2004). The human-
mouse GSTO1 orthologous pair has a low (r = 0.08), which is
not significantly different from 0 (P = 0.006; P = 0.98) (Supplementary Fig. 2). On
the other hand, the human and mouse GSTO2 orthologous genes,
which lack the alternative MER4A promoter, have significantly
correlated expression patterns (r = 0.76; P = 2.2e-6). These patterns of
expression divergence and conservation are consistent with
variation in expression introduced by the MER4A-TSS. In all, 37
out of 40 evaluated cases of human genes with ERV-TSS have
expression patterns that are not significantly correlated with
their mouse orthologs that lack the upstream ERV (Supplementary Fig. 3).

We further evaluated the potential regulatory effects of human
ERV-derived promoters by comparing the expression patterns
of all human genes with ERV promoters versus genes without
ERV promoters using the GNF2 data. Human genes with ERV-
TSS have greater tissue specificity than genes lacking ERV
promoters, consistent with a diversifying regulatory effect of
ERV-TSS (Supplementary Table 2). In particular, ERV-TSS
containing genes have anomalously high levels of expression,
on average, in brain and tests (Supplementary Figs 4 and 5).
A similar pattern of significantly elevated expression in brain
and tests was found for ERV CAGE tags (Supplementary Fig. 6).
Consistent with the brain-specific expression pattern of
ERV-TSS genes, GO functional analysis indicated that
these genes are enriched for metabolic and signaling processes
active in the brain (Supplementary Table 3 and Supplementary Fig. 7).

Our analysis revealed that retroviral sequences in the human
genome encode tens of thousands of active promoters; transcribed
ERV sequences correspond to 1.16% of the human genome sequence
and PET tags that capture transcripts initiated from
ERV's cover 22.4% of the genome. These data suggest that ERVs
may regulate human transcription on a large scale. However, it
is a formal possibility that many of the ERV derived promoters
identified here represent leaky transcription, i.e. noise,
which is not functionally significant. Definitive proof of biological
activity for individual ERV-TSS may have to await experimental
confirmation via knock-out data or promoter swapping. However,
it will soon be possible to validate ERV-TSS on a genome-
scale owing to the accumulation of high-throughput data from
tiling array experiments based on Chip-chip and/or chromatin
structure assays. Such data, which are being generated by the
ENCODe Project Consortium (2007), measure the distributions
of regulatory signatures across genomic sequence. The presence
and density of regulatory signals, such as transcription factor
binding sites and open or specifically modified chromatin,
have been shown to discriminate between biologically active
and artifactual TSS and thus could be used to validate
ERV-TSS.

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