Low conservation and species-specific evolution of alternative splicing in humans and mice: comparative genomics analysis using well-annotated full-length cDNAs

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ABSTRACT

Using full-length cDNA sequences, we compared alternative splicing (AS) in humans and mice. The alignment of the human and mouse genomes showed that 86% of 199 426 total exons in human AS variants were conserved in the mouse genome. Of the 20 392 total human AS variants, however, 59% consisted of all conserved exons. Comparing AS patterns between human and mouse transcripts revealed that only 431 transcripts from 189 loci were perfectly conserved AS variants. To exclude the possibility that the full-length human cDNAs used in the present study, especially those with retained introns, were cloning artefacts or prematurely spliced transcripts, we experimentally validated 34 such cases. Our results indicate that even retained-intron type transcripts are typically expressed in a highly controlled manner and interact with translating ribosomes. We found non-conserved AS exons to be predominantly outside the coding sequences (CDSs). This suggests that non-conserved exons in the CDSs of transcripts cause functional constraint. These findings should enhance our understanding of the relationship between AS and species specificity of human genes.

INTRODUCTION

Alternative splicing (AS), the recombination of exons to produce novel transcripts, is thought to contribute substantially to the functional complexity of human proteins. AS frequently produces diverse transcripts from a single genetic locus. The consequent exon changes alter the corresponding amino acid sequences and generate functionally divergent proteins. The ENCODE and GENCODE projects have found that about 60% of all loci encode AS isoforms, with an average of more than 5.4 isoforms per locus (1–3). In the ENCODE project, a product consisting of two exons of the DONSON gene and three exons of the ATP50 gene was cloned and sequenced by real-time reverse transcriptase polymerase chain reaction (RT-PCR) to generate several novel connected transcripts. Because many transcripts are derived from non-conserved sequences, however, the investigators in those projects emphasized the need for further studies exploring the neutrality of genome evolution. Several other reports have suggested that frequently occurring AS isoforms are not evolutionarily conserved and thus may be attributable to transcriptional noise or cloning artefacts.

Analyses of AS have relied on partially sequenced cDNA information (i.e. on expressed sequence tags, or ESTs). Although EST-based genome-wide approaches have identified thousands of instances of AS in human genetic transcripts (4,5), they do not provide information about the positions and combinations of AS exons in
full-length transcripts. This information is needed for
detailed analysis of the impact of AS on protein function
and evolutionary turnover. Furthermore, the standard
methods for estimating selective pressure by calculating
and comparing the rates of synonymous and non-
synonymous substitutions are not easily applied to par-
tially sequenced cDNA.

In the work reported here, we collected and analysed
instances of AS in the context of full-length cDNAs, which
are ideal resources with which to analyse AS because
information regarding the complete form of a particular
transcription unit (i.e. an isoform) allows us to determine
the relevance of AS to conserved protein functions. The
functional domains of proteins are often embedded over
a wide region of the protein sequences, so it is possible
that not every combination of AS exons is allowed. Full-
length cDNA-based approaches also facilitate better cov-
ereage at the 5’-ends of AS sequences than do EST-based
approaches. Our data are based on a large collection of
physical cDNA clones whose complete sequences have
been determined and therefore can thus be used to directly
validate the functions of certain genes. In a previous
study, we analyzed 56 419 full-length human cDNAs
whose sequences were checked by expert scientists and
by using computational methods specific to the full-
length cDNA annotation conferences H-Invitational
(H-Inv) (6) and H-Inv 2 (7). From that dataset, we created
a catalogue of 18 297 AS variants at 6877 loci (8).

Our AS catalogue is not the perfect tool with which to
understand the functional diversity of human genes.
Information about evolutionary conservation must be
added to our functional annotations if one is to use our
catalogue to ascertain which of the AS sequences in spe-
cific parts of proteins should be prioritized in future func-
tional analyses. Many studies have demonstrated that a
significant proportion of AS sequences are not conserved,
and those sequences can be assumed to have species-
specific biological roles. Also, it is suggested that not a
few genes seem to exist in a species-specific manner. We
therefore compared human AS variants with AS variants
in the mouse genome. We selected 431 conserved AS var-
ants at 189 loci for which full-length human and mouse
cDNAs were available. Interestingly, a significant number
of the AS variants that were not directly supported by full-
length mouse cDNAs, which contained non-conserved
exons, were translated to proteins. Here, we compare the
evolutionary conservation of AS transcripts in humans
and mice by using full-length cDNAs.

MATERIALS AND METHODS

Full-length cDNA sequences from humans and mice

We obtained 64 034 full-length human cDNAs sequenced
in four projects—Human Unidentified Gene-Encoded
Large Proteins (HUGE), Full-Length cDNA Japan
(FLJ), Munich Information Centre for Protein
Sequences (MIPS) and Mammalian Gene Collection
(MGC)—involving six institutions: Kazusa DNA
Research Institute, Tokyo University, Helix Research
Institute, German Cancer Research Centre, the United
States National Institutes of Health and the Chinese
National Human Genome Centre. These institutions pro-
vided full-length human cDNA clones that had been
annotated at the H-Inv and H-Inv 2 conferences (Supple-
mental Table 1A). We also obtained 118 775 full-length
mouse cDNAs that had been sequenced in the Functional
Annotation of Mouse (FANTOM) 3 and MGC projects
(Supplementary Table 1B). All sequences are registered in
release 66 of the DNA Data Bank of Japan (DDBJ) and
are freely available at http://www.ddbj.nig.ac.jp/.

Identification of human alternatively spliced variants

Human AS variants were identified as previously
described (8). To remove potential 5'- or 3'-truncated
cDNAs, we excluded sequences with 5'- or 3'-ends in the
second or downstream exons of other sequences. We
included cDNAs with 5'- or 3'-ends in the first or last
exon and which were therefore considered to vary in their
transcriptional initiation or termination sites. We
assumed that cDNAs with 5'-ends outside the exonic
regions of other sequences were not truncated forms of
known transcripts. For a detailed discussion of this
topic, see Kimura et al. (9). The resulting set of putative
full-length cDNAs was used to compare the genomic posi-
tions of each exon–intron boundary with that of other
transcripts from the same locus, with an allowance of
10 bp to remove potential sequencing or mapping errors
(Supplementary Table 2). If part of the cDNA exonic
sequence was in the first or last exon of another cDNA
intrinsic region, that sequence was considered a ’5'/3'-end’
AS variant. If part of the internal cDNA exonic sequence
was in the confirmed intrinsic region of another cDNA,
then that sequence was considered an ’internal’ AS vari-
ant. We then removed annotated genomic ‘rearrange-
ment’ genes, such as those encoding immunoglobulin
(Ig) and the T-cell receptor (TCR) and anomalously
highly polymorphic genes, such as those of the major his-
tocompatibility complex (MHC). If a group of AS vari-
ants contained two or more variants with the same
genomic structure, the one of median length was selected
as a representative of the group. If the number of variants
in the group is even, then the longer of the two near-
median-length variants was selected as a representative
of the group.

Functional annotation of human AS isoforms

We identified AS variants containing full-length open
reading frames (ORFs), which means the start codon
(ATG) encoded methionine and the final codon was
TAA, TAG or TGA. Full-length ORFs were defined as
coding sequences (CDSs). Of the 20 392 representative
human AS variants, 16 103 transcripts (i.e. 14 597 AS var-
ants) were determined to be isoforms with CDSs. Four
types of protein functions in these transcripts were anno-
tated: protein motifs, gene ontology (GO) terms, sub-cell-
ular localization signals and transmembrane domains.
Protein motifs and GO terms were identified with
InterProScan (10); sub-cellular localization signals were
predicted with WoLF PSORT (11) and TargetP (12) and
transmembrane domains were predicted with TMHMM
(13) and SOSUI (14) software. These annotation analyses of protein function were automatically executed by TACT (15), an integrated annotation tool for genome and transcriptome analyses. Further details regarding the functional annotation pipeline are available from H-InvDB (http://www.h-invitational.jp/) (16). The results of the computational identification and annotation analyses were visually inspected by members of the AS annotation team of H-Inv. Controversial annotations were flagged to identify possible errors.

Detection of retrotransposons and exonic splicing enhancers in human AS variants

RepeatMasker [A. F. A. Smit, R. Hubley and P. Green, RepeatMasker Open-3.0. 1996 to 2004 http://www.repeatmasker.org] was used to detect retrotransposons [long terminal repeats (LTRs), long interspersed nuclear elements (LINEs) and short interspersed nuclear elements (SINEs)] in the DNA sequences of human AS variants. Exonic splicing enhancers (ESEs) were detected in the DNA sequences of human AS variants by using 238 candidate hexamer sequences obtained from the RESCUE-ESE Web Server (17).

Alignment of the human and mouse genomes

Human and mouse genome sequences (versions hg18 and mm8) were downloaded from UCSC (http://genome.ucsc.edu/) and a modified pairwise genome alignment was constructed using BLASTZ (18) version 7, with parameter C = 2. Regions of redundant alignment were removed using reciprocal best hits. This method was originally used to construct two satellite databases of H-InvDB: the comparative genomics database G-compass (19) and the orthologue database Evola (20).

Genomic comparison of human AS variants and the mouse genome

The human–mouse genome alignment was used to compare the position of each human AS variant exon with that of the corresponding exon in mice. If the mapped coverage and identity of the overlapping regions surrounding the full-length or CDS in the exon exceeded threshold values (70% for coverage and 60% for identity), the region was considered 'genome-conserved'. If the full-length or CDS in the exon corresponded to a mouse counterpart transcript with coverage and identity equal to or greater than the threshold values, this region was considered 'transcript-conserved'. If the exon did not map to the genome alignment, or if the mapped coverage or identity was less than the threshold value, the exon was considered 'non-conserved' (Figure 1A).

The exon information was used to determine the conservation category for each transcript. If at least one non-conserved exon was identified in the transcript, the transcript was considered ‘non-conserved’. If the transcript had no non-conserved exons but contained genome-conserved exons, it was considered ‘genome-conserved’. If the transcript had only transcript-conserved exons, it was considered ‘transcript-conserved’ (Figure 1B). If all the exons in the human and mouse transcripts were transcript-conserved and the exon combination was also conserved, the corresponding transcripts were defined as equally spliced variants (ESVs). Finally, if different ESVs were identified at a particular human and mouse locus, the transcripts were considered ‘conserved AS’ (Figure 1C).

Figure 1. Comparative analysis of human and mouse AS sequences. (A) Categories of conserved exons. Full-length exons and coding regions of exons were included in the analysis, and the highest conservation level was selected. Boxes indicate exons; thin straight lines indicate introns. (B) Categories of conserved AS variants within the categorized exons in (A). Conservation levels were determined from the lowest conservation level of each transcript’s exons. (C) Equally spliced variants (ESVs) and conserved AS sequences represent higher categories of transcript-conserved variants. ESVs are transcript-conserved variants with similar combinations of exons in different species. Conserved AS sequences are two or more different ESV pairs at a particular locus in multiple species. Additional details are available in the Results section and the Materials and methods section.

Experimental validation of ‘retained-intron’ AS sequences

Polysomal fractions were prepared, using ~3 x 10^7 cells, as described elsewhere (21). Cell pellets were suspended in 1 ml of lysis buffer [20 mM Tris–HCl (pH 7.5), 10 mM NaCl, 3 mM MgCl_2, 0.04 M sucrose, 0.5% NP40 and 1 mM dithiothreitol] containing 100 units of RNase inhibitor. The pellets were then lysed by incubation on ice for 10 min. The nuclei and cell debris were removed by centrifugation at 1000g for 10 min at 4°C. The lysate was layered on top of 11 ml of a 15%/50% (w/v) sucrose gradient and centrifuged at 36000g in a Beckman SW41Ti
First-strand cDNA synthesis was performed using a 17-bp dT primer and SuperScript II as directed by the manufacturer. The resulting cDNAs were quantified and used in quantitative RT–PCR analysis. The results were normalized to the glyceraldehyde-3-phosphate dehydrogenase (GAPDH) PCR product. PCR-cycling parameters were 95°C for 2 min, 95°C for 10 min, followed by 35 cycles of 95°C for 30 s, 57°C for 1 min and 72°C for 1 min. The ABI PRISM HT7000 Sequence Detection System (Applied Biosystems) was used to detect the products. The primers used in the RT–PCR are described in Supplementary Table 3A. The sizes and integrity (the degree of smeared amplicons and primer dimers) of the final PCR products were confirmed by agarose gel electrophoresis.

RESULTS

Generation and characterization of evolutionarily conserved AS sequences

Evolutionarily conserved AS sequences were selected from the H-Inv and H-Inv 2 datasets. We examined the evolutionary conservation of exons in 20 392 AS variants of 64 034 human full-length cDNAs. The exons were defined as ‘conserved’, when the human genomic sequence to which the full-length or CDS regions of the exons were mapped could be aligned with the mouse genome with a coverage of >70% and an identity of >60%. Human–mouse alignments were performed using information available in the UCSC Genome Browser (versions hg18 and mm8). Of the 199 426 total exons, 171 547 (86%) were categorized as conserved (Table 1, panel A). Moreover, the exons in 12 096 transcripts of the 20 392 AS sequence variants were categorized as genome- or transcript-conserved (Table 1, panel B). On the other hand, in 944 human transcripts, all of the exons were not conserved. Therefore, the locus itself seemed not conserved. Although we did not analyze such putatively human-specific loci separately, they should be studied in greater detail (also see Supplementary Table 4).

The human transcripts were further examined to determine whether corresponding transcripts existed in mice. For this purpose, 118 775 full-length mouse cDNAs were collected and annotated during the FANTOM 3 and MGC projects to determine if any exons within the full-length mouse cDNAs mapped to corresponding genomic regions in humans. The conserved exons described above were separated into transcript-conserved and genome-conserved categories, if the coverage and identity of the mouse counterpart exon was more than the threshold value (Figure 1A). A transcript consisting of all transcript-conserved exons was deemed to be transcript-conserved (Figure 1B). The ESVs, in which all of the exons and their combinations were conserved, and the conserved AS sequences, in which both of the AS variants were ESVs, were more highly conserved categories of transcript-conserved variants (Figure 1C). These categories are explained in more detail in the Materials and methods section.

We then identified 4624 ESVs from 3570 loci and determined that only 431 AS variants from 189 loci were conserved (Table 1, panel C). Because these AS variants will undoubtedly prove interesting in future investigations, we confirmed their CDSs and looked for alternative splicing-mediated changes in the corresponding protein sequences and for the presence of protein motifs, GO terms, subcellular localization signals and transmembrane domains within the affected regions.

Examples of evolutionarily conserved AS sequences

An example of a conserved AS sequence is shown in Figure 2A. At the locus encoding the phosphoinositide-3-kinase (PI3-kinase) regulatory subunit, the 5’-exons of BC094795 are AS relative to another variant, BC030815. These human-transcript variants correspond to the mouse-transcript variants BC026146 and BC051106, respectively. All of the exons as well as their combinations and AS patterns are conserved in each of the human–mouse transcript pairs. Three AS variants (p85-α, p55-α and p50-α) were originally identified at the human

| Table 1. Genomic conservation of AS in humans and mice |
|-----------------------------------------------|
| Total | Non-conserved | Genome-conserved | Transcript-conserved |
|-------|---------------|------------------|----------------------|
| Panel A: exon-level conservation of human AS variants, compared with the mouse genome |
| All exons | 199 426 | 27 879 (14%) | 23 412 (12%) | 148 135 (74%) |
| AS exons | 49 842 | 12 196 (25%) | 9 064 (18%) | 28 582 (57%) |
| Panel B: transcript-level conservation of human AS variants, compared with the mouse genome and transcripts |
| AS variants | 20 392 | 8 296 (41%) | 4 410 (21%) | 7 686 (38%) |
| AS loci | 7 601 | 1 716 (23%) | 1 241 (16%) | 4 644 (61%) |
| Transcript-conserved total | | | | |
| Non-equally spliced variant | ESV | Conserved AS |
| Panel C: higher conservation categories for transcript-conserved AS sequences |
| AS variants | 7 686 | 2 631 (34%) | 4 624 (60%) | 431 (6%) |
| AS loci | 4 644 | 885 (19%) | 3 570 (77%) | 189 (4%) |
PI3-kinase locus in humans (22), with BC094795 corresponding to p85-α and BC030815 corresponding to p55-α. The functional domains specific to Rho GTPase-activating protein (RhoGAP) (IPR0088936) and Src homology-3 (SH3) (IPR001452) are embedded in the N-terminal region of p85-α, which extends from the N-terminal region of p55-α. The p85-α and p55-α variants appear to have diverged functionally to the extent that they relay signals from the insulin receptor substrate (IRS) proteins to PI3-kinase with different efficiencies (23). Because it seems natural that such a fundamental diversification would be evolutionarily conserved in both mice and humans, we were not surprised to identify these AS variants in mice. Indeed, they were categorized as ‘conserved’ in our study.

Figure 2B shows an example of a locus at which we identified previously unknown functional protein changes caused by AS. The protein products of this locus have been annotated as cysteinyl-tRNA synthetase (CARS). A glutathione S-transferase (GST) C-terminal-like (IPR010987) protein motif was identified in the BX647906 AS variant, but was absent from the BC002880 variant because of the presence of a cassette-type AS pattern in the second exon of BX647906. The C-terminal region of GST plays an essential role in substrate activation, and this motif occurs not only in GST but also in many types of aminoacyl-tRNA synthetases (aaRSs) (24). Some types of mammalian aaRSs interact with translational elongation factor (EF)-1 via this GST C-terminal domain, and this interaction is thought to facilitate the vectorial transfer of aminoacylated tRNAs. Thus, the functional roles of the newly identified AS variants BX647906 and BC002880 may involve the delivery of the cognate tRNA from CARS to the EF complex, thereby controlling translation efficiency in the cell under specific circumstances (25).

Figure 2. Conserved AS sequences, exemplified by (A) PI3-kinase regulatory subunit and (B) CARS. Constitutive introns are shortened here and additional details are available in the Results section. Boxes indicate exons. Filled regions within boxes indicate CDSs (green), protein motifs (red) and untranslated regions (yellow). The ESVs shared by humans and mice are indicated by arrows.

### Table 2. Comparison of AS features in total and conserved AS loci

| Panel A: AS patterns | Total | Conserved |
|---------------------|-------|-----------|
| Total               | 7601  | 189       |
| Cassette (Skipped exon) | 3584 (35%) | 66 (42%) |
| Internal acceptor (Alternative 3′ splice) | 1988 (19%) | 30 (19%) |
| Internal donor (Alternative 5′ splice) | 1990 (20%) | 33 (21%) |
| Mutually exclusive | 237 (2%) | 4 (2%) |
| Retained intron    | 2477 (24%) | 26 (16%) |

| Panel B: Effects of AS on protein function | Total | Conserved |
|-------------------------------------------|-------|-----------|
| Total                                     | 7601  | 189       |
| Average difference in CDS length          | 87 aa | 52 aa     |
| Total effects of AS on function            | 3395 (45%) | 125 (66%) |
| Protein-motif alterations                  | 2423  | 86        |
| GO alterations                             | 1078  | 50        |
| Sub-cellular localization changes          | 2305  | 75        |
| Transmembrane domain changes               | 444   | 16        |
We selected 14 retained-intron AS variant pairs from the transcript-conserved population, 15 from the genome-conserved population, and 5 from the non-conserved population, which were manually annotated as ‘highly likely protein coding’ sequences. In the transcript-conserved population all of the exons were conserved at the transcriptional level (i.e. there were corresponding mouse cDNAs for both variants). In the genome-conserved population, there were no corresponding mouse transcripts but the corresponding genomic sequences for all the related exons were conserved. In the non-conserved population, at least one of the related exons was not conserved. We determined the expression patterns of the selected variants in 20 types of human tissues by using semi-quantitative real-time RT–PCR.

As a control, we performed the same series of experiments on adjacent exons separated by moderately long introns. We observed no evidence of immature splicing in any of the controls. PCR products corresponding to the retained introns were observed in at least one tissue for 14 of the 14 transcript-conserved variants, for three of the 15 genome-conserved variants and for three of the five non-conserved variants. A wide variety of expression patterns was observed. For example, we observed ubiquitous expression of both type of some AS variants (Figure 3A, upper panel), whereas we observed the mutually exclusive expression of other AS variants in a tissue-preferred manner (Figure 3A, lower panel). These results suggest that a significant population of cDNAs, even those resulting from retained-intron AS variants, were derived from real transcripts. The overall expression patterns and mutual dependence of the AS variants appear to be controlled.

We further investigated whether transcripts containing retained-intron AS sequences were translated into proteins. We did this by recovering RNAs from actively transcribing ribosomal fractions (i.e. polysomal fractions) in a human promyelocytic leukaemia cell line, HL60, with sucrose density-gradient purification (Figure 3B). RNAs purified from fractions 7–10 were analysed with real-time RT–PCR. RT–PCR of total RNA revealed that 14 of the transcript-conserved retained introns that were expressed at least one tissue were expressed in HL60 cells. For nine of these, clear bands of the appropriate size were produced by RT–PCR of the polysome fractions (Figure 3C and D and Supplementary Table 3B).

### Table 3. GO terms and protein motifs frequently observed at conserved AS loci

| GO ID          | Definition                     | Number of conserved AS loci | Total number of AS loci | P  |
|----------------|--------------------------------|-----------------------------|-------------------------|----|
| GO:0003677     | DNA binding                    | 17                          | 333                     | 0.0085* |
| GO:0004601     | Peroxidase activity            | 3                           | 14                      | 0.0077* |
| GO:00060979    | Response to oxidative stress   | 3                           | 14                      | 0.0077* |

P-values were calculated using Fisher’s exact test. They indicate the significance of the difference between the ratios to the conserved AS loci (189) and the total AS loci (7601). *P < 0.01.

(IPR004827) motif, is also significantly more common in the conserved AS loci (Table 3, panel B). In eukaryotes, the bZIP transcription factor is responsible for initiating cellular responses to ultraviolet (UV) damage and osmotic stress (28). Because the switch of these functional motifs is essential to basic cellular functions, the corresponding AS sequences should also be conserved between species.

These results suggest that the functions of AS sequences differ between conserved AS variants, which constitute the core dataset of the AS sequences and have conserved diversification of gene functions, and other AS variants. In other words, a distinct subset of protein motifs appears to be responsible for the functional diversity of conserved and species-specific gene functions. Classification of the AS variants is necessary to further address this issue.

### Experimental validation of retained introns

After selecting and manually annotating the AS variants, we sought to understand why there was such a large population of non-conserved AS variants. One possibility is that these non-conserved variants are artefacts derived from aberrant cDNA cloning (e.g. cDNAs produced from incompletely spliced mRNAs or genomic DNA contaminants). Sequence information is insufficient to distinguish truly functional retained-intron type AS sequences (which are less common among conserved AS sequences) from the rest of the population.

We selected 14 retained-intron AS variant pairs from the transcript-conserved population, 15 from the genome-conserved population, and 5 from the non-conserved population, which were manually annotated as ‘highly likely protein coding’ sequences. In the transcript-conserved population all of the exons were conserved at the transcriptional level (i.e. there were corresponding mouse cDNAs for both variants). In the genome-conserved population, there were no corresponding mouse transcripts but the corresponding genomic sequences for all the related exons were conserved. In the non-conserved population, at least one of the related exons was not conserved. We determined the expression patterns of the selected variants in 20 types of human tissues by using semi-quantitative real-time RT–PCR.

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### Characterization of non-conserved exons

To further characterize the non-conserved exons, we first compared the characteristic features of the non-conserved...
We also observed that non-conserved exons were more frequently related to retrotransposons (Table 4). Even in cases where non-conserved AS exons overlap with protein motifs, the categories of overlapping motifs differ strikingly from those motifs overlapping conserved AS sequences. The most frequently observed protein motif in non-conserved AS exons was the KRAB box (IPR001909) (Supplementary Table 6) (30). This observation is consistent with the results of a previous study reporting that this motif, which is a well-known protein interaction domain, was significantly enriched in the sites of alternative splicing (31). The KRAB-type zinc finger is thought to control the transcription of the sex-determining region Y (SRY/Sry) gene and is therefore essential for the determination and differentiation of the testis (32). The fact that KRAB motifs tend to overlap with non-conserved AS exons may underscore the evolutionary uniqueness of the human transcriptional regulatory network in the reproductive system. A few protein motifs, including the GAGE motif (IPR008625), nuclear-pore-complex-interacting motif (IPR009443), phospholipase A2 active site motif (IPR013090) and the T-complex 11 motif (IPR008862), were also identified only in non-conserved AS exons (Supplementary Table 6). The GAGE motif is specific to humans and the nuclear-pore-complex-interacting motif is specific to primates (33,34). We further examined the expression patterns of these genes having GAGE, nuclear-pore-complex-interacting, phospholipase A2, active site and T-complex 11 motifs by using our expression profiling database H-ANGEL (Human Anatomical Gene Expression Library) (35). We found that the genes containing the GAGE and T-complex motifs were preferentially expressed in testis, that the expression pattern of genes containing the nuclear-pore-complex-interacting motif was ubiquitous and that there was no data on the expression pattern of genes containing the phospholipase-A2-active-site motif (Figure 4). Although the functions of GAGE-motif-containing genes are potentially very interesting, all that is known about that them is that they are primate specific. We were unable to investigate the AS of the GAGE motifs in this study because their parent genes are not found in mice (also see Supplementary Table 6). The parent genes for the T-complex-11 motifs, in contrast, are found in both humans and mice. Furthermore, human and mouse TCP11, which encodes the receptor for fertilization-promoting peptide (FPP), is reported to play an important role in sperm function (36). In any case, it is interesting that the GAGE and T-complex-11 motifs, which are enriched in the non-conserved AS exons, were associated with a species-specific factor involved in the reproductive system.

Our results also show that non-conserved exons are more frequently related to retrotransposons (Table 4). This observation is not surprising because human retrotransposons are primate specific (Alu) and mouse retrotransposons are rodent specific (37). Exonic-splicing enhancers (ESEs) are ubiquitously scattered in human AS variants (Table 4), so we could not observe the relation between ESE conservation and splicing (38). We found no clear differences between the non-conserved and conserved

exons with those of the conserved exons. Our conserved exon data set consisted of the genome level and above, and our results showed that the majority of non-conserved exons are located at the 5′-end of the transcript and appear to function as AS exons (Supplementary Table 5). Consistent with our findings, recent studies have found that alternative 5′-exons derived from alternative promoters are abundant in humans and are often not evolutionarily conserved (29). In contrast, the coding regions of non-conserved exons overlap less frequently with AS exons than with constitutively spliced (CS) exons (Table 4). It is likely that previous studies reported overall higher frequencies for non-conserved exons among AS exons because the number of non-conserved exons is greatest at the 5′-ends of non-coding exons.

**Figure 3.** Experimental validation of AS human transcripts by using the retained-intron pattern. (A) RNA expression of retained-intron AS sequences in 20 human tissues (1, adrenal gland; 2, bone marrow; 3, brain, cerebellum; 4, brain, whole; 5, fetal brain; 6, fetal liver; 7, heart; 8, kidney; 9, liver; 10, lung, whole; 11, placenta; 12, prostate; 13, salivary gland; 14, skeletal muscle; 15, testis; 16, thymus; 17, thyroid gland; 18, trachea; 19, uterus and 20, spinal cord). The upper panel exemplifies ‘ubiquitous’ retained-intron AS sequences and the lower panel exemplifies ‘tissue-preferred’ retained-intron AS sequences. (B) RT–PCR analysis using polysomal fractions isolated from the human promyelocytic leukaemia cell line HL60. (C) RNA expression of retained-intron AS sequences mixed with translating ribosome fractions (i.e. polysome fractions) from the HL60 cell line. (D) Number of expressed transcripts in each conserved category.
Table 4. Relationship between conservation and splicing in human alternatively spliced variant exons

|                | Total         | CDS-related | Protein-motif-related | Retrotransposon-related | Exonic-splicing-enhancer-related |
|----------------|---------------|-------------|-----------------------|------------------------|----------------------------------|
| C/CS exons     | 133901        | 95583 (71%) | 27805 (21%)           | 2523 (2%)              | 130104 (97%)                     |
| C/AS exons     | 37646         | 20805 (55%) | 6192 (16%)            | 2308 (6%)              | 35702 (95%)                      |
| NC/CS exons    | 15683         | 7030 (45%)  | 1898 (12%)            | 2549 (16%)             | 14961 (95%)                      |
| NC/AS exons    | 12196         | 3516 (29%)  | 812 (7%)              | 4544 (37%)             | 11701 (96%)                      |
| All exons      | 99426         | 126934 (64%)| 36707 (18%)           | 11924 (6%)             | 192468 (97%)                     |

C, conserved; NC, non-conserved.

DISCUSSION

We performed a large-scale comparative study of the AS variants in human transcripts, a study based on completely sequenced and intensively annotated full-length human and mouse cDNAs. Our dataset of AS full-length cDNAs is unique in that all combinations of AS exons have been defined as single entities in the complete form of the transcript. To the best of our knowledge, this is the largest available dataset of its kind. Although some of the AS variants identified here, especially those having an easily detectable internal AS pattern like a cassette, have been previously identified with EST-based and microarray-based approaches and are in the RefSeq and Ensembl databases, the previous identification of those variants was based on the interpretation of fragmented partial cDNA sequences and computational gene predictions. Indeed, 12710 of the 13705 cassette AS exons identified in this study (93%) had been identified in more than one EST-based study (Supplementary Table 7A). On the other hand, only 11% of the cassettes AS exons we identified in this study are in the RefSeq database and only 38% of them are in the Ensembl database (Supplementary Table 7B). We think that one of the major barriers to these AS exons being included in the representative gene models has been a lack of precise annotations, which are now available in our dataset. Our manually and computationally inspected data allowed us to examine the genome-wide features of AS sequences in a far more reliable manner than was previously possible, and our findings will enhance our understanding of the complex biological nature of AS sequences. A very interesting paper based on our AS database revealing the relation between transcriptional start sites (TSSs) and AS has already been published (39).

This study provides the first glimpse of the evolutionary turnover of AS sequences in full-length transcripts. Throughout evolution, the genomes of higher eukaryotes have expanded and mutations have been introduced by environmental factors (e.g. ultraviolet radiation) or internal factors (e.g. transposons). Mammals and other higher eukaryotes are thought to have accumulated numerous novel AS variants, and many AS sequences identified in this study appear to have occurred in a human-specific manner. By integrating information about evolutionary conservation with functional annotations, we have determined that non-conserved exons appear to have unexpectedly been excluded from the CDSs (Table 4). We had hypothesized that the CDSs would be more likely to contain non-conserved AS exons than conserved ones because the selective pressure against their biological functions is weak. We however found evidence of selective pressure against non-conserved AS exons in the CDS. Thus, it is likely that exons are easily generated by alternative splicing after speciation but are more likely to produce malfunctioning proteins when they occur in the CDS. It is also interesting that most AS variants are produced in the brain and testis (40). Because these tissues contain large numbers of neurons and sperm cells that could be functionally served as mutual substitutions or because variability itself may be important in these tissues, the occasional malfunction of individual proteins might be relatively well tolerated there. Thus, cells in the brain and testis might take advantage of the opportunity to experiment with AS variants, and this experimentation might act as a driving force in evolution and speed the process of speciation.

We shall soon overcome the current problems associated with the use of full-length cDNA data, the greatest of which—the limited number of sequences available—prevents us from analysing all AS variants. Researchers are already initiating large-scale projects by, using the next generation of DNA sequencers (e.g. Illumina 1G and SOLiD) to sequence previously isolated and partially...
sequenced cDNA clones. The same type of full-length cDNA data now available for mice and men will soon become available for other mammals. Further enrichment of the expression information will also allow us to precisely analyse in what tissue the species-specific AS sequences are utilized. The current presumption that the products of AS are found most frequently in brain and testis should be tested by non-EST-based sampling-bias-free analysis. Cataloguing full-length cDNAs and creating related databases are the first steps toward understanding how AS variants contribute to the functional and evolutionary diversification of gene functions.

SUPPLEMENTARY DATA
Supplementary Data are available at NAR Online.

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