Monotreme-specific conserved putative proteins derived from retroviral reverse transcriptase

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Abstract

Endogenous retroviruses (ERVs) have played an essential role in the evolution of mammals. ERV-derived genes are reported in the therians, many of which are involved in placental development; however, the contribution of the ERV-derived genes in monotremes, which are oviparous mammals, remains to be uncovered. Here, we conducted a comprehensive search for possible ERV-derived genes in platypus and echidna genomes and identified three reverse transcriptase-like genes named RTOM1, RTOM2, and RTOM3 clustered in the GRIP2 intron. Comparative genomic analyses revealed that RTOM1, RTOM2, and RTOM3 are strongly conserved and are under purifying selection between these species. These could be generated by tandem duplications before the divergence of platypus and echidna. All RTOM transcripts were specifically expressed in the testis, possibly suggesting their physiological importance. This is the first study reporting monotreme-specific de novo gene candidates derived from ERVs, which provides new insights into the unique evolution of monotremes.

Key words: endogenous retrovirus; monotreme; reverse transcriptase; virus-derived gene.

Introduction

Endogenous retroviruses (ERVs) are remnants of retroviral genomes found in the host genomes. ERVs are retroviruses that infected the host germline cells and were integrated into the host genome (Johnson 2019). Young ERVs retain their viral open reading frames (ORFs) but gradually lose their intact ORFs due to the accumulation of mutations. However, proteins expressed from ERVs sometimes evolve as functional genes in the host (Ueda et al. 2020). A typical example is the syncytin genes, ERV-derived fusogenic genes, which are expressed in the human placenta (Blaise et al. 2003; Blond et al. 2000; Mi et al. 2000) and are essentially required for mouse placenta formation (Dupressoir et al. 2009, 2011). Syncytin genes have been independently acquired from different ERVs in different mammalian lineages, which is a representative example of the convergent evolution (Imakawa, Nakagawa, and Miyazawa 2015). In addition, other ERV-derived genes that do not show fusogenic activity have also been found to be expressed in the placenta. For example, HEMO encoding a secreted envelope protein (Heidmann et al. 2017) as well as gagV1 and pre-gagV1 genes (Boso et al. 2021) are highly expressed in the human placenta. Restriction factors against exogenous retroviruses are another example of viral gene co-option. For example, gag-derived Fv1 (Best et al. 1996) and env-derived Fv4 (Ikeda and Sugimura 1989) inhibit retroviral infection in mice.

Despite these contributions to the evolution of therians, it is still unclear whether ERV-derived genes are co-opted in monotremes (egg-laying mammals).

Here, we attempted to determine whether there are ERV-derived genes specific to monotremes. Comparative studies for the detection of ERV-derived genes have been conducted in mammalian genomes, including the platypus (Nakagawa and Takahashi 2016; Wang and Han 2020). However, for monotremes, only the genome sequence of one species, the platypus, was available (OANA5), the quality of which was limited (Warren et al. 2008). Recently, high-quality monotreme genomes of platypus (mOrnAna1.p.v1) and echidna (mTacAcu1.pri) were determined using long-read sequencing technology (Zhou et al. 2021). Taking advantage of these genome sequences, we conducted comparative analyses and detected three novel ERV-derived genes specific to the monotreme lineage.

Results and discussion

To comprehensively search for ERV-derived genes in the genomes of monotremes, we extracted ORFs from the genomes of platypus and echidna. The amino acid sequences obtained by the virtual translation of these ORFs were used as queries for the sequence search. We used the hidden Markov model (HMM) of retroviral genes in the Gypsy Database 2.0 (GyDB) (Llorens et al. 2011) as
Figure 1. Identification of RTOM1, RTOM2, and RTOM3. (A) Schematic representation of the in silico screening for conserved ERV-derived genes in platypus and echidna. (B) Genomic context of RTOM1, RTOM2, and RTOM3. The thin arrows indicate ORFs above 100 amino acid length. The nucleotide sequences of RTOM ORFs are available in Supplementary File 2.
genomes (mammals: 26 out of 510 genomes, birds: 29 out of 556 genomes, and reptiles: eight out of seventy-nine genomes) (Supplementary Table S4), however, their query cover rates were low (up to 5.2 per cent) except for hits to the RTOM ORFs themselves. The tBLASTn search identified up to thousands of hits for each genome (Supplementary Table S4). This is because the amino acid sequences of the RTOM ORFs are similar to the RT region of other ERVs. We examined the proximity of these hits to the GRIP2 gene and found no hits considered to be orthologs of the RTOM ORFs (see the ‘Materials and Methods’ section). Therefore, we conclude that the RTOM ORFs are monotreme-specific.

We found that, in the platypus genome, there are computationally annotated RefSeq genes containing the RTOM ORFs (Fig. 2A). RTOM1, RTOM2, and RTOM3 genes of platypus contain two introns in the 5′ UTR, and the entire RTOM ORFs are expressed as mRNA excluding a second splicing variant of RTOM3 that partially lost its ORF (Fig. 2A). In echidna, RTOM2 and RTOM3 gene structures were annotated in the RefSeq transcripts; however, RTOM1 was not annotated. By conducting transcriptome assemblies of RNA-seq data of echidna tissues (Supplementary Table S5), we reconstructed all RTOM transcripts including the RTOM1 (Fig. 2B, Supplementary Fig. S4). We also found a chimeric transcript of RTOM2 and RTOM3, which was transcribed from the transcription start site of RTOM2, but its CDS is RTOM3 (Supplementary Fig. S4). Except for this chimeric transcript, all echidna RTOM transcripts have two introns in the 5′ UTR, which was similar to those of platypus. We then constructed a multiple alignment of the seven amino acid sequences of platypus and echidna RTOMs, including two splicing variants of platypus RTOM3 (Fig. 2C). The amino acid sequence of RTOM2 lacks a region shared by RTOM1 and RTOM3, but the C-terminal region was conserved among the amino acid sequences of RTOMs without insertion or deletion (Fig. 2C). To investigate the tissue-specific expression of the RTOM genes, we analyzed the RNA-seq data of platypus and echidna (Supplementary Table S5). In platypus, RTOM1, RTOM2, and RTOM3 were commonly highly expressed in the testis (Fig. 2D). GRIP2 was expressed not only in the testis but also in the brain, and its expression level was lower than that of the RTOM genes. We further investigated the mapped reads using Interactive Genome Viewer (Thorvaldsdottir, Robinson, and Mesirov 2013) (Supplementary Fig. S5) and found that RTOM3 showed a splicing variant with an intron in the coding region, as shown in the RefSeq transcript. In echidna, we found that all RTOM transcripts were specifically expressed in the testis, similar to platypus. Expression of GRIP2 in echidna testis was also relatively low (Fig. 2E). This suggests that the RTOM genes are more actively transcribed than GRIP2, or the RTOM transcripts are more stable than the GRIP2 transcripts in the testis. Given the higher expression level of RTOM2 in both platypus and echidna, this gene may play a central role in the RTOM genes. It is still possible that the relative expression levels of three genes may change according to other tissues and developmental stages that were not examined in this study. In addition, since this study did not present the evidence of the translation of the RTOM transcripts, whether putative RTOM proteins are involved in testicular function needs to be verified in the future.

To obtain insights into the viral origin of the RTOM genes, we performed a BLASTp search for the amino acid sequence of platypus RTOM1 against the NCBI virus database. We found that retroviral Pol proteins from various distinct lineages, namely gammaretrovirus, deltaretrovirus, epsilonretrovirus, and spumaretrovirus, are similar to the amino acid sequence of RTOM1 (BLASTp: E-value <1E−20). In all hits, the retroviral Pol proteins showed high similarity to the latter half of RTOM1 (approximately 370-607aa) (Fig. 3A). A domain search against the Pfam database (Mistry et al. 2021) in the HMMER web service (Finn, Clements, and Eddy 2011) revealed that the latter half of RTOM1 and RTOM3 contain RT domains (Supplementary Fig. S6). A phylogenetic tree was constructed from the RT regions of the amino acid sequences of RTOMs and the retroviral Pol proteins (Fig. 3B). The amino acid sequences of RTOMs appear to be more related to class III retroviruses, including spumaviruses or spumavirus-related MuERV-L (Llorens et al. 2009). The tree topology of the RTOMs strongly suggests that all the RTOM genes were formed before the divergence of platypus and echidna. Together with the self-alignment of genomic sequences (Supplementary Fig. S2), three RTOM genes were generated by tandem gene duplications before the divergence of platypus and echidna (Fig. 3C). In the non-RT region of the amino acid sequences of RTOM1 (approximately 1–369aa), no significant hits for retroviruses were obtained (Fig. 3A). We performed a BLASTp search for all non-redundant proteins in the GenBank database for the non-RT region of the amino acid sequences of RTOM1, however, no similar proteins were found except for the putative proteins of RTOM2 and RTOM3 (E-value <0.05). This suggests that the non-RT region was derived from a non-retroviral sequence or was derived from the retroviral gene that has accumulated too many mutations to be aligned with retroviruses. Considering the structural divergence of the non-RT region, such as deletion of RTOM2 and splicing variant of platypus RTOM3 (Fig. 2C), the RT region is a core domain of the putative RTOM proteins, and the non-RT region may provide functional modifications specific to each putative RTOM protein.

During the 187-million-year history of diverging from monotremes, therians have acquired many ERV genes and evolved their unique features, especially the placenta (Inakawa and Nakagawa 2017). Our work revealed that monotremes also domesticated ERV genes that emerged and were conserved more than 55 million years ago, the divergence time of platypus and echidna (Zhou et al. 2021). Although the translation of RTOM genes was not confirmed in this study, the calculation of nonsynonymous and synonymous nucleotide substitution frequencies of RTOM1, RTOM2, and RTOM3 shows that their amino acid sequences are under purifying selection, strongly suggesting that they physiologically function as proteins (Fig. 3D). We found that the RT domain of all putative RTOM proteins lacked the three catalytic carboxylates of aspartic acids (Supplementary Fig. S7). These amino acid residues are highly conserved among all retroviruses, and the replacement of these amino acids results in a complete loss of the RT activity (Larder et al. 1987; Sarafianos et al. 2009). Therefore, the putative RTOM proteins may have different functions from those of reverse transcription. To the best of our knowledge, there are no retroviral genes in which only the RT domain is opted in vertebrates (Naville et al. 2016). Although this study has a limitation that the expression and function of the putative RTOM proteins have not been fully validated due to difficulties in obtaining tissues, the future functional elucidation of RTOM1, RTOM2, and RTOM3 will provide us with new aspects of ERV-derived genes functioning in mammals.

Materials and methods
Identification of conserved ERV genes
The platypus genome (mOrnAna1.p.v1, GCF_004115215.1) and the echidna genome (mTAcAcu1.pri, GCF_015852505.1) were used for the ERV gene screening. The 240-nt ORF flanked by stop codons

K. Kitao et al. | 3
Figure 2. Expression of RTOM1, RTOM2, and RTOM3. (A) Schematic representation of the RefSeq transcripts of the RTOM genes in platypus. (B) Schematic representation of the reconstructed RTOM1 transcript and RefSeq transcripts of the RTOM2 and RTOM3 genes in echidna. (C) Multiple alignment of the amino acid sequences of RTOMs. The amino acid sequence of echidna RTOM1 was obtained from the genomic ORF ‘RTOM3A_platypus’ and ‘RTOM3B_platypus’ are isoforms derived from ‘XM_039910408.1’ and ‘XM_029051290.2,’ respectively. The regions showing similarity to the HMM of spumaretrovirus RT domain in GyDB are indicated in red. (D, E) Tissue-specific expression of RTOM genes and GRIP2 in (D) platypus and (E) echidna. Normalized expression levels are presented as transcript per million (TPM).

was retrieved using the getorf program in the European Molecular Biology Open Software Suite (Rice, Longden, and Bleasby 2000). For HMM-based sequence search, hmmscan was used (expected threshold: 1E−5) in HMMER3 v3.2.2 (Eddy 2011). ORFs were clustered using CD-HIT v4.8.1 (Li and Godzik 2006) with 50 per cent amino acid identity. The sequence search for platypus
ORFs against echidna ORFs was conducted using BLASTp v2.10.0+ with an e-value < 1E−50 (Camacho et al. 2009). By examining the distribution of the bitscore of the BLASTp search, we extracted ORF pairs that showed high similarity between the two species (Supplementary Fig. S1A). Then, we examined the RefSeq annotation and identified the genes to which the ORFs belonged.

To examine the presence of homologous sequences beyond the monotreme lineage, the genes that were not described in the human RefSeq genes were subjected to deep homology searches. We performed BLASTn and tBLASTn v2.10.0+ with e-values < 1E−5 against all genomes of mammals, birds, and reptiles downloaded on 22 January 2022 (Supplementary Table S4). The query cover rate for each hit was calculated as [alignment length/query length]. To examine whether the amino acid sequences obtained by tBLASTn are the ortholog of the RTOM genes, we investigated the proximity of these hits to the GRIP2 gene as follows. First, we extracted hits to the amino acid sequences of RTOMs with a query cover of at least 60 percent. Second, to obtain the genomic position of GRIP2 on each genome, we performed BLAT v0.35 (Kent 2002) using the amino acid sequence of the human GRIP2 (NP_001073892.3) against the 1,145 genomes. We considered the hit with the highest score in each genome as the GRIP2 position. Finally, we compared the genomic position of the hits to RTOM and GRIP2 and confirmed that all of them were located on different contigs or were located far enough from each other (the closest pair of hits to RTOM and GRIP2 in the same contig is 7.7 Mb apart from each other).
To validate this approach, we performed similar analyses on the genomes of humans (GRCh38.p13) and marmosets (Callithrix jacchus_cj1700_1.1) (Supplementary Fig. S1B and C). They diverged 43 million years ago (Perelman et al. 2011). We successfully identified known ERV-derived genes such as PEG10 (Ono et al. 2001), RTL1/PEG11 (Charlier et al. 2001), ASPRV1 (Matsui et al. 2011), NYNLIN/CGIN1 (Marco and Marín 2009), ERV-1 and 2 (Kjeldbjerg et al. 2008), and ERVMER34-1/HEMO (Heidmann et al. 2017) (Supplementary Table S6). This suggests that our method is sensitive enough to identify ERV ORFs conserved in platypus and echidna.

Expression analyses
RNA-seq data of platypus (twenty samples from six tissues) (Marin et al. 2017) and echidna (eleven samples from seven tissues) (Zhou et al. 2021) were used (Supplementary Table S5). Low-quality reads were trimmed and filtered using fastp v0.19.5 with default options (Chen et al. 2018). The filtered reads were mapped to each reference genome using HISAT2 v2.1.0 with default option (Pertea et al. 2016). Based on the eleven RNA-seq sequencing data mapped on the echidna genome, we obtained the echidna RTOM1 transcript by conducting transcriptome assembly using Stringtie2 v2.1.6 with ‘-merge’ option (Kovaka et al. 2019). We have added the coordinates of the echidna RTOM1 transcript (Supplementary File 1) to the RefSeq gene coordinates. We then calculated the expression levels for twenty platypus and eleven echidna RNA-seq samples using the Stringtie2 program with default options (Kovaka et al. 2021). To extract uniquely-mapped reads from a given aligner-generated SAM file, we collected reads containing the ‘NH:i:1’ flag indicating that they were uniquely mapped into the genome.

Phylogenetic analyses
Representative retroviral Pol amino acid sequences were retrieved from the GyDB collection (https://gydb.org/index.php/Alignment?alignment=POL_retroviridae_Biology_Direct_4_41_2009&format=txt) (Llorens et al. 2009). A multiple alignment was generated using MAFFT v7 487 (Katoh and Standley 2013), and poorly aligned regions were removed using trimAl v1.4 rev15 (Capella-Gutiérrez, Silla-Martínez, and Gabaldón 2009). A phylogenetic tree was constructed using IQ-TREE2 v2.0.8 (Minh et al. 2020) with 1,000 replicates of ultrafast-bootstrap (Hoang et al. 2018). The tree was visualized using FigTree v1.4.4 (http://tree.bio.ed.ac.uk/software/figtree/).

Supplementary data
Supplementary data are available at Virus Evolution Journal online.

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