

Minireview

3'UTR Diversity: Expanding Repertoire of RNA Alterations in Human mRNAs

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Genomic information stored in the DNA is transcribed to the mRNA and translated to proteins. The 3' untranslated regions (3'UTRs) of the mRNA serve pivotal roles in post-transcriptional gene expression, regulating mRNA stability, translation, and localization. Similar to DNA mutations producing aberrant proteins, RNA alterations expand the transcriptome landscape and change the cellular proteome. Recent global analyses reveal that many genes express various forms of altered RNAs, including 3'UTR length variants. Alternative polyadenylation and alternative splicing are involved in diversifying 3'UTRs, which could act as a hidden layer of eukaryotic gene expression control. In this review, we summarize the functions and regulations of 3'UTRs and elaborate on the generation and functional consequences of 3'UTR diversity. Given that dynamic 3'UTR length control contributes to phenotypic complexity, dysregulated 3'UTR diversity might be relevant to disease development, including cancers. Thus, 3'UTR diversity in cancer could open exciting new research areas and provide avenues for novel cancer therapeutics.

Keywords: 3'UTR diversity, alternative polyadenylation, alternative splicing, cancer, RNA alterations, transcriptome

INTRODUCTION

In the nucleus, eukaryotic mRNA is synthesized from a gene in a 5' to 3' direction and processed into mature transcripts by 5'-capping, splicing, and 3'-end formation. Genomic information is translated to proteins once mature mRNAs are exported to the cytoplasm. 5' and 3' sequences flanking the coding regions are not translated, thus named as untranslated regions (UTRs). What are the roles of UTRs in regulating gene expression? 5'UTRs are the leading sites for ribosome assembly for mRNA translation. In contrast, 3'UTRs play various roles in the post-transcriptional control of gene expressions, including stability, translation, and subcellular localization of mRNAs (Mayr, 2019). We will begin by reviewing why 3'UTR diversity is important, especially in human mRNAs. This review focuses only on the length of 3'UTRs, but not on their sequences and modification, as 3'UTR diversity for simplicity.

Evolutional features of human mRNA 3'UTRs

In general, the noncoding part of the genome has increased in size and complexity during evolution (Pesole et al., 2002). The average length of 5'UTRs is similar in metazoans; whereas, the 3'UTR is significantly longer in humans compared to other species (Mayr, 2016; Pesole et al., 2002; Sood et al., 2006; Wang et al., 2019) (Fig. 1). 3'-end sequencing also revealed that at least half of human genes could generate alternative 3'UTR isoforms from the same genes, demonstrating high diversity in 3'UTR in the human transcriptome (Derti et

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al., 2012; Hoque et al., 2013).

3'UTRs contain cis-regulatory elements recognized by trans-acting factors. Thus, changes in 3'UTR length could alter a number of regulatory elements (Ji et al 2009). In fact, many genes have alternative 3'UTR or contain internal introns which can switch 3'UTR length. Therefore, 3'UTR length changes could affect gene expression by fine-tuning and reprogramming the mRNA regulatory landscape in human cells (Navarro et al., 2021).

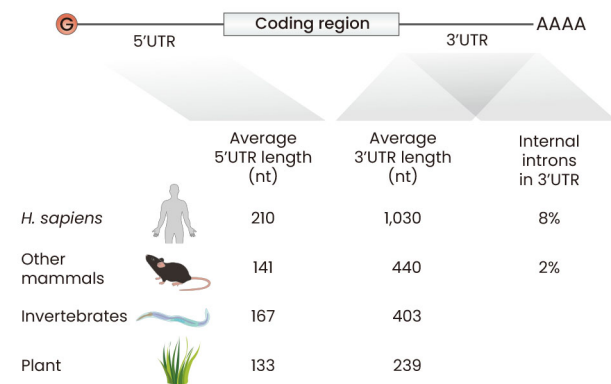


Fig. 1. Length and internal introns in untranslated region (UTR) during evolution. The average length of the 5'UTR and 3'UTR of mRNA among different organisms. The 5'UTR is typically shorter compared to the 3'UTR. In *Homo sapiens*, the average length of the 3'UTR is about 1030 nucleotides (nt), which is longer than the average length of the 3'UTR in other species. Percentage of genes containing internal introns is also shown in *H. sapiens* (8%) and other mammals (2%).

FUNCTIONS OF 3'UTRs

3'UTRs contain cis-regulatory elements recognized by trans-acting factors. 3'UTR-mediated mRNA metabolism controls at the steps of the stability, translation, and localization of mRNA (Fig. 2). 3'UTRs form ribonucleoprotein (RNP) with many RNA binding proteins (RBPs). So we will describe representative mRNA 3'UTRs, corresponding main RBPs, and their relevant functions as follows (Table 1).

Stability control

Most mRNAs undergo stability regulation through specific sequences and structures in 3'UTRs, such as AU-rich elements (AREs), GU-rich elements, CA-rich elements, and miRNA-binding sequences. Some proteins bind to AREs and recruit the degradation machinery to ARE-containing mRNAs. For example, tristetraprolin (TTP) binds AREs within the 3'UTRs of mRNAs, such as *tumor necrosis factor (TNF)*, *prostaglandin-endoperoxide synthase 2 (PTGS2/cyclooxygenase-2, [COX-2])*, *vascular endothelial growth factor A (VEGFA)*, *interleukin 10 (IL10)*, and *immediate early response 3 (IER3)*, and promote mRNA degradation (Essafi-Benkhadir et al., 2007; Lai et al., 1999; 2006; Sawaoka et al., 2003).

In contrast, Hu-antigen R (HuR) protein binds AREs and stabilizes numerous mRNAs, increasing the corresponding protein level (Brennan and Steitz, 2001). HuR stabilizes mRNAs promoting tumor growth and cell survival, such as *VEGFA*, *COX-2*, *TNF*, *CTNNB1/β-catenin*, and *interferon regulatory factor (IRF9)* (Dean et al., 2001; Rothamel et al., 2021; Sengupta et al., 2003; Steinman, 2007; Thiele et al., 2006). It also stabilizes mRNAs involved in cell cycle regulation, such as *cyclin dependent kinase inhibitor 1A (CDKN1A/p21)* and *cyclin (A2, B1, D1, and E1)* (Giles et al., 2003; Guo and Hartley, 2006; Wang et al., 2000). In addition, mRNAs encoding pro-survival proteins such as *FOS (c-fos)* and *B-cell CLL/lymphoma 2 (Bcl-2)* are stabilized by HuR (Ishimaru et al., 2009; Wang

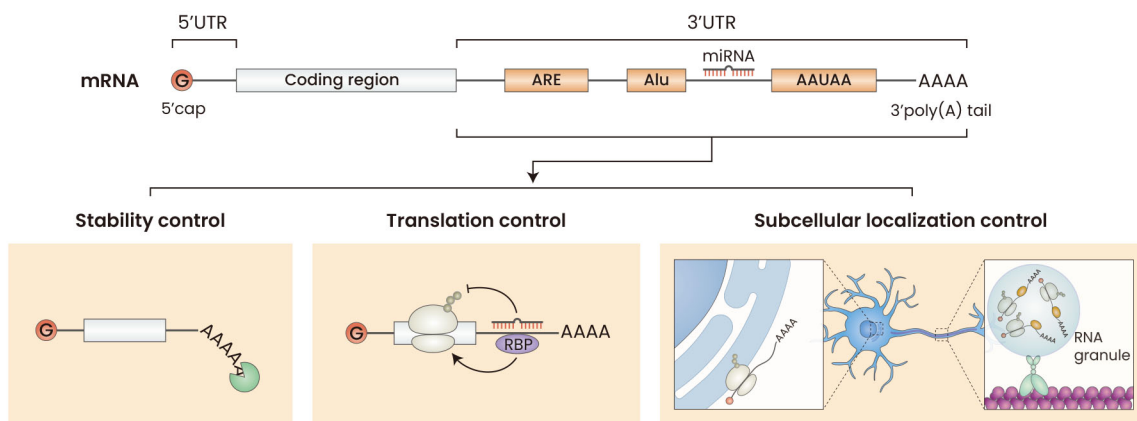


Fig. 2. Roles of 3' untranslated region (3'UTR) in post-transcriptional regulation. The diagram illustrates a typical mRNA structure consisting of 5'UTR, coding region, and 3'UTR. The 3'UTR is located following the stop codon and before the poly(A) tail. It includes cis-regulatory elements like AU-rich elements (AREs), Alu elements and microRNA (miRNA) targets, and polyadenylation sites. The 3'UTRs play important roles in post-transcriptional gene regulation of mRNA stability, translation, and localization.

Table 1. List of representative mRNAs with 3'UTR-mediated control

Transcript	RNA-binding proteins	Control	Function	Reference
<i>Hunchback</i>	Pumilio Nanos Brat	Translation repression	Embryonic axis formation in <i>Drosophila</i>	(Kuersten and Goodwin, 2003)
<i>UCP1</i>	CPEB2	Translation activation	Maintain body temperature	(Chen et al., 2018)
<i>CaMK2A</i>	Mub	Localization translation	Control learning and memory in neuron	(Broix et al., 2021; Chen et al., 2022)
<i>ACTB</i>	IGF2BP1 PAT1	Stabilization local-translation	Constituent of the cytoskeleton for structural support and movement in neuron	(Wu et al., 2020)
<i>Oskar</i>	Bruno Hrp48	RNP granule formation Translation	Embryonic patterning and germline formation in <i>Drosophila</i>	(Bose et al., 2022; Jamboret al., 2014)
<i>DMPK</i>	CUG-BP2	mRNA export Translation	Maintenance of the myelin sheath	(Amack and Mahadevan, 2001; Taneja et al., 1995)
<i>n-myc</i>	ELAV like RNA binding protein family	ARE-mediated mRNA stabilization	Cellular growth in neuroblastoma	(Chagnovich and Cohn, 1996; Chagnovich et al., 1996)
<i>c-fos</i>	SRSF3	Translation repression	Regulator of cell cycle and senescence	(Kim et al., 2022)
<i>PZ1^{IMAF1}</i>	HuR	mRNA stabilization	Cellular proliferation and tumorigenesis	(Kim et al., 2012; Lee and Jeong, 2006)
<i>COX-2</i>	b-catenin	Translation activation		

et al., 2001). These data suggest that cis-regulatory elements in 3'UTR participate in the context- and condition-dependent mRNA stability control by binding to the combination of different sets of RBPs.

Translation control

3'UTRs also contain cis-regulatory elements related to translation control. During the development of *Drosophila* embryo, *Hunchback* mRNA 3'UTR forms an RNP complex with Pumilio, Nanos, and Brat protein, leading to translation repression for proper development in the posterior region of the embryo (Kuersten and Goodwin, 2003). Likewise, many developmental genes are expressed by the translation regulation mediated by 3'UTR-RNP complexes.

Different lengths of 3'UTR contribute to the translation efficiency control due to the presence or the absence of cis-regulatory elements. In the case of Uncoupling protein 1 (UCP1), the long 3'UTR isoform is predominant and contains the binding sites for the translation regulator, cytoplasmic polyadenylation element-binding protein 2 (CPEB2) (Chen et al., 2018). CPEB2 is required for associating the long 3'UTR isoform for the low-level translation of *UCP1* mRNA in steady-state conditions, and translation upregulation in response to cold exposure or adrenaline (Chen et al., 2018). These data suggest the role of 3'UTRs for translation regulation by binding to many RBPs forming RNP complexes. 3'UTR-RNP-mediated translational regulation is also linked to mRNA localization, as described below.

Localization control

The subcellular localization of mRNA is investigated mostly in neurons (Andreassi and Riccio, 2009). Localized mRNA is translated in specific subcellular locations and allows the precise production of proteins in spatiotemporal patterns in response to neural activity. For example, calcium/calmodulin-dependent protein kinase II alpha (CaMK2A) regulates calcium signaling in the nervous system as a critical player in activity-dependent behavioral plasticity (Bae and Miura, 2020). In *Drosophila*, the 3'UTR of *CaMK2A* mRNA contains regulatory information necessary for local translation in presynaptic terminals (Broix et al., 2021). In mammals, the resting concentration of CaMK2A should be maintained extremely high, reaching up to 2% of the total protein in the hippocampus. This enrichment is achieved through the enhanced axoplasmic translation of *CaMK2A* mRNA, which is mediated by the RBP Mub and a 23-base Mub-recognition element in 3'UTR (Chen et al., 2022).

Actin beta (*ACTB*/ β -*actin*) mRNA is another example of 3'UTR-mediated mRNA transport. 3'UTR of *ACTB* mRNA associated with the zipcode-binding protein 1 (ZBP1), also known as insulin-like growth factor 2 mRNA-binding protein (IGF2BP) (Wu et al., 2020). This association facilitates the transport of *ACTB* mRNAs to the specific locations in neurons, such as dendrites, in response to brain-derived neurotrophic factor (BDNF) (Wu et al., 2020). Overall, the 3'UTR of the *ACTB* mRNA is crucially involved in its transport and localization in neurons, which is essential for the proper development and function of the nervous system.

The control of mRNA localization is also investigated in

Drosophila embryo. In the case of *oskar* mRNA, the 67-nt stem-loop structure in 3' UTRs is essential for transporting the mRNA from the nurse cells into the oocyte. This transport depends on the action of dynein and kinesin motor proteins, which use the oocyte entry signal to guide the mRNA to its proper location within the embryo (Bose et al., 2022; Jambor et al., 2014). Therefore, 3' UTRs of an mRNA molecule are critically involved in determining its subcellular localization and manipulating its translation. In this way, 3' UTRs can regulate the spatiotemporal expression of a gene and control when and where a protein is produced.

Multi-step control

During the lifecycle of mRNAs, some RBPs bind to 3' UTRs and affect multiple steps of post-transcriptional regulation (Moore, 2005). Such multifunctional RBPs associate with 3' UTRs of many mRNAs involved in critical downstream events. Quaking (QK) is an RBP with a STAR domain involved in compact myelin formation (Vernet and Artzt, 1997). QK binds to a specific sequence in the 3' UTRs of several mRNAs that encode proteins related to myelin formation. The QK response element (QRE) is "ACUAAAY" in 3' UTRs of mRNAs, such as *microtubule associated protein 1B* (*MAP1B*), cyclin dependent kinase inhibitor 1B (*CDKN1B/p27kip*), and heterogeneous nuclear ribonucleoprotein A1 (*hnRNP A1*) mRNAs (Larocque et al., 2005; Li et al., 2000; Zearfoss et al., 2011; Zhao et al., 2006).

Some proteins are newly found RBPs implicated in multiple 3' UTR-mediated control steps. For example, β -catenin is a Wnt-activated transcription factor and an adhesion protein. In addition, it also associates with ARE in 3' UTRs of *COX-2* mRNAs as an RBP in colon cancer cells. *COX-2* protein expression is regulated by mRNA stability and translation control by 3' UTR-binding β -catenin in the cytoplasm (Lee and Jeong, 2006). Furthermore, the interaction between β -catenin and *COX-2* mRNA assembles a tertiary complex with the HuR protein, vital for the upregulation of *COX-2* protein levels and colon cancer progression (Kim et al., 2012). Considering that *COX-2* protein is the enzyme critical for synthesizing prostaglandins, a high level of β -catenin RBP might have significant pathological effects on cancer cells.

Another example is serine/arginine-rich splicing factor 3 (SRSF3). SRSF3 is a member of the serin/arginine-rich (SR) family proteins, acting in AS mainly in the nucleus. In addition, the shuttling activity of SRSF3 is linked to AS regulation, export, and translation in the cytoplasm (Park and Jeong, 2016). SRSF3 interacts with the 3' UTR of *p21^{cip1/waf1}* (*CDKN1A*) mRNA and represses translation in the cytoplasm (Kim et al., 2022).

These multifunctional RBPs are located in both the nucleus and cytoplasm and form 3' UTR-mediated RNP to control the multiple steps of mRNA metabolism. Further research is needed to fully understand the multiple roles of RBPs in 3' UTR-mediated mRNA regulation.

REGULATION OF 3' UTR FUNCTIONS

Considering the important roles of 3' UTRs in post-transcriptional gene expression, 3' UTR-mediated control should be

regulated by many factors, including miRNA, RBPs and RNA granules as follows.

By the interplay of miRNA and RBPs

3' UTR functions are regulated by the crosstalk of miRNAs and RBPs, and cis-regulatory elements within 3' UTRs. miRNAs are small noncoding RNA molecules that are crucially involved in regulating gene expression by binding to complementary sequences in the 3' UTRs of the target mRNA. This binding can lead to the destabilization of mRNAs or the inhibition of their translation (Baek et al., 2008; Friedman et al., 2009). For example, *VEGFA* mRNA 3' UTR contains several regulatory elements, including CA-rich elements and miRNA-binding sequences. In normal conditions, miR-297 and miR-299 bind to the 3' UTR of the *VEGFA* mRNA leading to reduced expression of *VEGFA* proteins (Matoulkova et al., 2012). However, under hypoxic conditions, the silencing activity of miR-297 and miR-299 on *VEGFA* mRNA is inhibited by the binding of hnRNPL to CA-rich elements in 3' UTRs, allowing mRNAs to be continuously translated into proteins (Jafarifar et al., 2011). These data suggest that the interplay of miRNAs and RBPs are important for the condition-dependent regulation of 3' UTR activities.

The control of 3' UTR length also could affect miRNA-binding sites and leads to functional consequences (Nam et al., 2014). Thus, the dynamic regulation of 3' UTR length could play a significant role in development because it contributes to the gene expression patterns observed in different cell types and tissues. Likely, distinct sets of genes are differentially regulated by the interplay of miRNA and RBPs in a context- and condition-dependent manner (Hoffman et al., 2016).

By forming RNA granules

RNA granules in the cytoplasm are emerging as essential players in mRNA localization and translation regulation (Moore, 2005; Tian et al., 2020). RNA granules are specialized structures in assembled RNAs with RNP particles. Several RNA granules include transport RNPs, stress granules, and processing bodies (P-bodies) (Kiebler and Bassell, 2006). This compartmentalization creates cellular asymmetries, may enhance biological reactions, and promote molecular interactions required for cell growth and development (Tian et al., 2020). How RNAs assemble RNA granules? 3' UTRs within mRNAs could be key components in forming RNA granules. For example, the cytoplasmic polyadenylation element (CPE) in 3' UTR and its binding protein CPEB facilitate mRNA transport to dendrites in rat hippocampal neurons (Huang et al., 2003). 3' UTR-RNP granules are formed on *CaMK2A*, *MAP2*, and *β -catenin* mRNAs with CPEB, maskin, and eIF4E, for transporting corresponding mRNAs (Huang et al., 2003; Kundel et al., 2009; Rook et al., 2000).

Technical advances enhance cellular RNA granule visualization. *In situ* hybridization (ISH) demonstrated the *bicoid* and *oskar* RNP granules with Staufen double-strand binding protein in *Drosophila* oocytes (Huang et al., 2003; Kiebler and Bassell, 2006; St Johnston, 2005). Moreover, the improvement of *in situ* RNA visualization methods, such as single-molecular fluorescent ISH (smFISH) and RNAscope[®] made it

possible to observe *oskar* RNP granules with scaffold protein Bruno and Hrp48 (Bae and Miura, 2020; Bose et al., 2022; de Planell-Saguer et al., 2010). Further enhancement of mRNA visualization techniques will be required to elucidate and visualize the 3'UTR-dependent RNA granule localization.

GENERATING 3'UTR LENGTH DIVERSITY

3'UTR diversity can be caused by various ways, such as genetic variations, RNA modification, alternative cleavage, alternative polyadenylation (APA), and alternative splicing (AS). These could diversify the transcriptome and contribute to epigenetic gene regulation. This section focuses on the length-diversifying mechanism, APA, and AS (Fig. 3). Databases with different 3'UTR lengths formed by APA and AS are also summarized in Table 2.

APA in 3'UTR

Cleavage and polyadenylation is a critical step in the maturation of 3'UTR ends of most eukaryotic mRNAs (Mitschka and Mayr, 2022). The polyadenylation process begins the recognition of specific sequences in the mRNA molecule by the polyadenylation machinery in the nucleus. The polyadenylation machinery consists of several factors, including cleavage factor I, cleavage and polyadenylation specificity factor, cleavage factor II, and cleavage stimulation factor. These complexes bind to specific sequences in mRNA 3'UTRs and facilitate the cleavage at a specific site downstream of the AAUAAA motif. This cleavage is aided by cleavage factor polyribonucle-

otide kinase subunit 1, followed by adding a polyadenylate tail to the mRNA molecule by the enzyme poly(A) polymerase alpha (Tian and Manley, 2013). A strong polyadenylation signal at the 3'UTR end is important for the efficient cleavage and polyadenylation of mRNAs. However, many genes also have additional polyadenylation signals at their 3'UTRs in eukaryotes, and the usage of these signals can be regulated through APA (Fig. 3A).

APA is a widespread phenomenon in metazoan protein-coding transcripts (70%-79% of mammalian genes). In humans, >70% of genes have more than one polyadenylation site in their 3'UTRs, and approximately 50% have three or more. By contrast, in the mouse liver, >60% of expressed genes had multiple polyadenylation signals in their 3'UTRs (Tian and Manley, 2013). This can result in mRNAs with different 3'UTR ends and different regulatory and functional properties (Mitschka and Mayr, 2022). 3'UTR shortening was proposed to increase mRNA stability by reducing the accessibility of mRNA degradation mechanisms, such as RBP- or miRNA-based degradation. Conversely, 3'UTR lengthening may increase the accessibility of miRNAs, decreasing mRNA stability and translation.

In addition to regulating mRNA stability APA can play a role in mRNA-protein interactions and protein localization via alternative 3'UTRs. For example, the APA of *CD47* mRNA 3'UTR can affect the interaction with different protein complexes and thereby direct *CD47* localization to either the plasma membrane (long 3'UTRs) or endoplasmic reticulum (short 3'UTRs) in various cancers (Berkovits and Mayr, 2015;

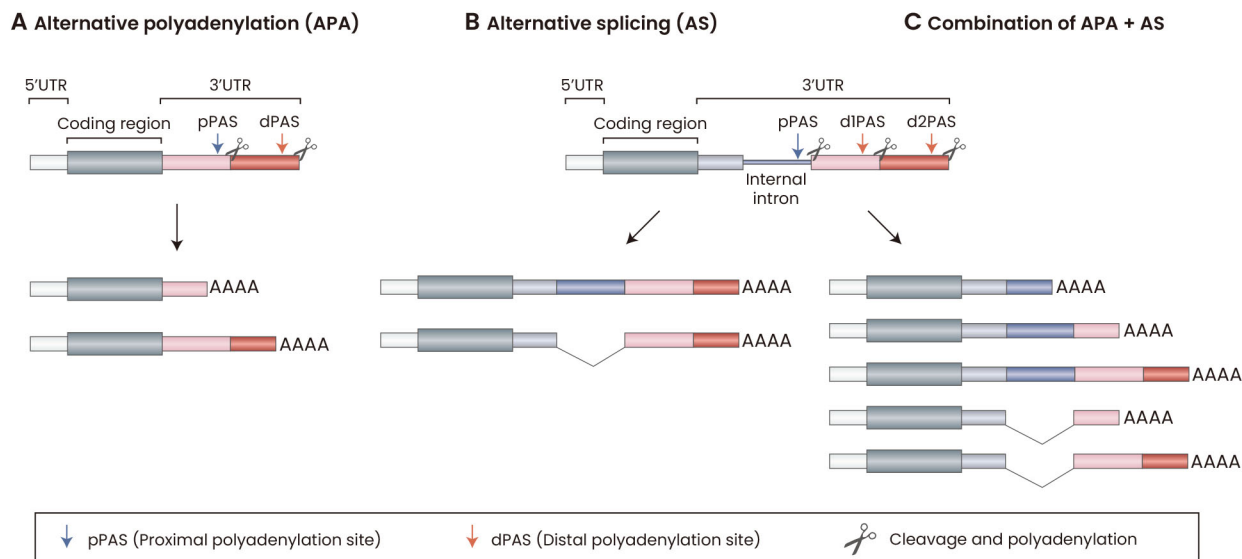


Fig. 3. Regulatory events generating 3' untranslated region (3'UTR) length diversity. (A) Alternative polyadenylation (APA) in 3'UTR. The generation of 3'UTR length diversity using different cleavage and polyadenylation sites: proximal polyadenylation site (pPAS) and distal polyadenylation site (dPAS). The selection of pPAS over dPAS, or vice versa, can result in different 3'UTR lengths. If the pPAS is selected, the 3'UTR will be shorter (shown in the pink box); the dPAS is selected, the 3'UTR will be longer (shown in the pink box and red box). (B) Alternative splicing (AS) within 3'UTR. The generation of 3'UTR length diversity using the internal intron in 3'UTR (shown in the blue line). Inclusion or skipping of this intron can affect the length of the 3'UTR. If the intron is included, 3'UTR will contain an intronic sequence (shown in the blue box). (C) Combination of APA and AS events in 3'UTR. More 3'UTR variants can be generated.

Table 2. List of Database related to 3' UTR divers

Tool name	Features	URL	Reference
Poly(A) sites databases			
PolyASite 2.0	APA atlas made from 3' end sequencing data	https://polyasite.unibas.ch	(Herrmann et al., 2020)
PolyA_DB 3	APA atlas built from ~1.2 billion 3' end deep sequencing reads	https://exon.apps.wistar.org/PolyA_DB/v3/	(Wang et al., 2018)
scAPAdb	APA atlas at single-cell resolution from publications and Genomics website	http://www.bmibig.cn/scAPAdb/	(Zhu et al., 2022)
TREND-DB	APA database constructed from bulk RNA-seq data of potential APA regulators	http://shiny.imbei.uni-mainz.de:3838/trend-db	(Marini et al., 2021)
3' UTR alternative splicing databases			
RNA structure databases	3' UTR splicing events database from RNA-seq data of TCGA and GTEx	https://www.cbrc.kaust.edu.sa/spur/home/	(Chan et al., 2022b)
RNA structure databases			
RNAfold	Prediction tool of secondary structures for single-stranded RNA or DNA	http://rna.tbi.univie.ac.at/cgi-bin/RNAWebSuite/RNAfold.cgi	(Lorenz et al., 2011)

Ma and Mayr, 2018). More studies are required to understand this interesting phenomenon by 3' UTR-directed protein localization.

AS in 3' UTRs

AS can also alter the 3' UTR length if 3' UTR contains an internal intron (Fig. 3B). It will increase the length, heterogeneity, and functional diversity of 3' UTRs (Bicknell et al., 2012; Chan et al., 2022b). The combination of APA and AS could further increase mRNA isoform variability resulting in the generation of mRNA transcripts containing various forms (Fig. 3C). The splicing of introns located within 3' UTRs was first described in *heterogenous nuclear ribonucleoprotein D (hnRNP/D/AUF1)* (Wilson et al., 1999). This study reported that splicing of the intron in 3' UTR regulates AUF1 expression, so it may play a significant role in gene expression regulation.

Another study demonstrated that 3' UTRs of *integrin subunit alpha 3 (ITGA3)* mRNA interacts with muscleblind like splicing regulator 2 (MBNL2) in human lung carcinoma cells (Adereth et al., 2005; Pascual et al., 2006). These splicing factors regulate the AS of *ITGA3* 3' UTR, in which produces distinct 3' UTR variants with different functions. The authors also showed the *ITGA3* mRNA localized to adhesion plaques, specialized structures involved in cell adhesion and signaling. Moreover, in the gene coding for an *activity regulated cytoskeleton associated protein (Arc)*, AS generates isoforms with distinct 3' UTRs. The AS of *Arc* 3' UTR is induced by long-term potentiation learning and memory (Paolantoni et al., 2018).

The 3' UTR of the *β-catenin* gene *CTNNB1* is also an interesting example. Three 3' UTR variants are expressed following the AS of an intron located within 3' UTRs (Chan et al., 2022b; Thiele et al., 2006). The expression of variant 3' UTRs in *β-catenin* mRNA was reported in peripheral blood mononuclear leukocytes (PBMCs). This suggests that different isoforms of *β-catenin* mRNAs may play different roles in regulating gene expression in PBMCs (Thiele et al., 2006). Recently, a pan-cancer analysis revealed pervasive upregulation of 3' UTR splicing driving tumorigenesis with *β-catenin* as a top-dysregulated 3' UTR (Chan et al., 2022b). Thus, 3' UTR

diversity could not only contribute to the overall diversity of the transcriptome and proteome but also drive cancer development and progression.

3' UTR-MEDIATED DISEASES

Transcriptome analyses of human cancer databases revealed that a large percentage of genes in cancers have RNA alterations (PCAWG Transcriptome Core Group et al., 2020). RNA alterations are mostly in protein-coding regions, but 3' UTR diversity also attribute to RNA alterations. Because APA is widespread and alters the regulatory potential of 3' UTRs, APA dysregulation plays a significant role in cancers (Kahles et al., 2018). Global APA within 3' UTRs is involved in the proliferation and metastasis of cancer cells and tumor tissues (Hoque et al., 2013; Mayr and Bartel, 2009). APA dysregulation is exemplified in *Cyclin D1* overexpression because of 3' UTR shortening (Mayr and Bartel, 2009; Wiestner et al., 2007). This 3' UTR shortening leads to the loss of binding sites of complementary miRNA to escape repression and potentially contribute to oncogene activation (Yang and Nam, 2020).

Based on genome and transcriptome databases, AS events in cancer tissues are estimated to be 20% higher on average than in normal tissues (Kahles et al., 2018). Recently, a pan-cancer analysis revealed that pervasive upregulation of 3' UTR splicing drives tumorigenesis (Chan et al., 2022b). In hepatocellular carcinoma, aberrant 3' UTR isoforms of *β-catenin* mRNA escape non-sense-mediated mRNA decay and promote cell proliferation and migration (Chan et al., 2022b). Thus, 3' UTR diversity could not only contribute to the overall complexity of the transcriptome and proteome but also drive cancer development and progression (Chan et al., 2022a).

As in cancer disorders of neuronal plasticity and learning, 3' UTRs appear to be a pathological hotspot (Conne et al., 2000). Because several neuronal mRNAs depend on their 3' UTRs for appropriate subcellular targeting or translational control, perturbations in 3' UTR-mediated functions deserve attention. In addition, alterations in the secondary structure of

3'UTRs are related to the pathology of a certain disease (Reamon-Buettner et al., 2007). Mutations in the 3'UTR of *GATA binding protein 4* (*GATA4*) mRNA alter its secondary structure and may contribute to the development of congenital heart disease (Reamon-Buettner et al., 2007). Therefore, further studies could explain the mechanism of the 3'UTR-mediated disease development (Conne et al., 2000).

CONCLUSIONS AND PERSPECTIVES

As genomic and transcriptomic data accumulated, RNA alterations emerged as important features of cancer development (PCAWG Transcriptome Core Group et al., 2020). Among many RNA alterations, 3'UTR variants received little attention in comparison with those in protein-coding regions (Bicknell et al., 2012). However, a widespread shortening of 3'UTR lengths in cancers and causal UTR variants in human diseases demonstrated the important roles of 3'UTRs in disease development (Griesemer et al., 2021; Mayr and Bartel, 2009). 3'UTR alterations can contribute to the diversification of the transcriptome and protein functions, impacting numerous biological processes.

3'UTR activities are context- and condition-dependent, dynamically affecting gene expression programs by miRNAs, RBPs, and RNA granule formation. Thus, finding rules and factors generating and modulating 3'UTR variants, such as 3'-end processing machinery, 3'UTR-binding proteome and UTR-mediated RNP granules, might be important. As functional alterations in 3'UTRs are found in various diseases, a systemic search for "3'UTR-mediated diseases" will extend the horizons in the diagnosis and treatment of these diseases (Conne et al., 2000). Specifically, functional cis-regulatory elements in 3'UTRs in pathogenic genes, interaction with disease-associated RBPs, and their pathogenic conditions should be understood to elucidate the mechanism and their consequent phenotypic changes in diseases. It could also open exciting new research areas, answering the question of what enables the biological complexity in humans during evolution.

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AUTHOR CONTRIBUTIONS

S.J. conceived the idea, wrote the manuscript, and secured funding. D.W. performed experiments and wrote the manuscript.

CONFLICT OF INTEREST

The authors have no potential conflicts of interest to disclose.

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