**Introduction**

N^6^-methyladenosine (m^6^A) is one of the common forms of RNA modifications, which is found to occur in RNAs of many organisms or cell lines (1-9). The distribution of m^6^A on mRNAs are not random, it is reported that the methylated adenosine is located in GRAC, RRACH or DRACH motif (R = A or G, H = A, C or T, D = A, G or T). In mammals, one out of one thousand adenosines in total RNAs are N^6^-methylated (5,6,8,10). Over ten thousand m^6^A modification sites have been identified in mRNAs (4-6,8,11,12), which are enriched in long exons, 3'UTRs and around stop codons. The m^6^A methylation events are being regulated by different factors and meanwhile regulating the downstream biological processes. The dynamic regulation of m^6^A modification is accomplished by the m^6^A writers (methyltransferase complex) (13,14) and erasers (15,16), and the downstream effects of m^6^A are exerted with the help of readers (5,6,11,17) or by m^6^A itself (1,4).

The writer complex is mainly composed of METTL3, WTAP and METTL14. METTL3 is the core catalytic

**Background:** N^6^-methyladenosine (m^6^A) is one of the common forms of RNA modifications. METTL3 is the essential factor that has methyltransferase activity. One important role of m^6^A is to regulate translation of mRNAs via reader YTHDF1. However, whether m^6^A could indirectly affect the translation of mRNA through other approaches remains unclear.

**Methods:** We retrieved the m^6^A genes in HeLa cells generated by a previous study. In the METTL3 or YTHDF1 knock-down libraries, we examined the global changes in mRNA splicing as well as translation efficiency (TE).

**Results:** In METTL3-KD cells, the differential splicing (DS) genes are enriched in m^6^A modified genes. The DS events are relatively enriched in 5'UTR of mRNAs. The 105 genes with DS events in 5'UTR alter their TE more strongly than the genes with DS events in other regions (CDS/3'UTR/intron). Furthermore, the splicing pattern of 98 out of those 105 genes are unaffected by reader YTHDF1. Importantly, we did not observe significant TE changes for these 98 genes when YTHDF1 was knocked down.

**Conclusions:** In HeLa cells, for a small set of genes, m^6^A could modulate the translation of modified mRNAs through affecting the splicing patterns. These indirect effects are independent of the direct regulation by reader proteins as we have verified using YTHDF1-KD data. This pattern is likely caused by the gain or loss of cis-elements in 5'UTRs that determine the translation of host genes. Our work extended our knowledge about the translation regulation by m^6^A.

**Keywords:** N^6^-methyladenosine (m^6^A); translation efficiency (TE); differential splicing (DS); cis-element; indirect
enzyme and METTL14 acts as a structural role (18). WTAP is found to stabilize the dimerized METTL3 and METTL14. Other proteins like RBM15 and KIAA1429 were also found in the writer complex (19). Apart from the preference in RRACH motifs, based on the fact that the methylation process is co-transcriptional (20), recent studies revealed that METTL14 interacts with histone modifications and promote site-specific methylation on pre-mRNAs (21). The reversibility of methylation indicates the presence of eraser proteins. Two erasers FTO and ALKBH5 are found in mammals. The lack of FTO leads to abnormal growth, development, mobility and metabolism (22-24). The mutated ALKBH5 would affect spermatogenesis, and also cause cancer (25,26).

The reader proteins of m^6^A modification are important for executing the biological functions of m^6^A. In mammals, m^6^A readers mainly include YTHDF1/2/3 and YTHDC1/2. Phenotypical and functional studies discovered that the readers are important for stem cell development and cancer immunity (27-29). Meanwhile, the mechanistic studies revealed profound molecular details of how m^6^A sites interact with reader proteins. Binding of YTHDF2 on m^6^A transcripts affects the stability of the mRNAs (5), and YTHDF1 recruits the translation initiation factors (particularly eIF3) to m^6^A modified mRNAs to enhance the translation efficiency (TE) of these genes (6). Meanwhile, m^6^A is also able to modulate mRNA translation independent of reader proteins. Under heat stress, m^6^A on 5'UTR of Hsp70 mRNAs acts as IRES (internal ribosome entrance site) to facilitate the translation initiation (by eIF3) of Hsp70 (9,30).

The cases of YTHDF1 (6) and IRES (9,30) are both related to the initiation factor(s) eIF3 that could directly determine the translation initiation. We wonder whether m^6^A could fine-tune the translation process through other indirect approaches. Interestingly, it is reported that m^6^A modification could alter the splicing patterns of mRNAs and this change even affects the sex determination in Drosophila (1,3,4). This also indicates the case that m^6^A modification could exert its function without the need of reader proteins. This phenomenon reminds us that m^6^A might be able to modulate translation through other indirect ways. It is established that many cis-elements in the 5'UTR are important for determining the TE of downstream mRNA, such as sequence motifs for scanning, structures and non-canonical reading frames. If the splicing changes caused by m^6^A could lead to the gain or loss of these decisive cis-elements in 5'UTRs, this would eventually affect the translation of host genes.

To test our hypothesis, we retrieved the m^6^A genes in HeLa cells. In the METTL3 knock-down libraries (6), we examined the global changes in mRNA splicing patterns as well as TE. The differential splicing (DS) genes are enriched in m^6^A modified genes and the DS events are relatively enriched in 5'UTRs. The 105 genes with DS events in 5'UTR alter their TE more strongly than the genes with DS events in other regions (CDS/3'UTR/intron). Furthermore, the splicing pattern of 98 out of those 105 genes are unaffected by reader YTHDF1. Importantly, we did not observe significant TE changes for these 98 genes when YTHDF1 was knocked down.

Our results demonstrate that m^6^A could modulate the translation of mRNAs through affecting the splicing patterns, at least for a small set of genes. These indirect effects are independent of the direct regulation by reader proteins. Our work extended our knowledge about the translation regulation by m^6^A.

Methods

Next generation sequencing (NGS) data

The NGS (next generation sequencing) data in normal HeLa cells or HeLa cells with si-METTL3/si-YTHDF1 were downloaded from a previous study (6). The NGS data contain mRNA-Seq and Ribo-Seq (ribosome profiling followed by deep sequencing) (31) that enable us to define the TE of each gene. Those adenosine sites in GRAC motif (if located in m^6^A peaks) were systematically recognized as m^6^A modification sites. Note that throughout this manuscript, the “si-” and “KD” mean the same. They are used to avoid monotonic expressions.

Assigning the m^6^A sites to the human genes

We downloaded the hg19 human genome (UCSC). In determining whether an m^6^A site is located in the 5'UTR/ CDS/3'UTR/intron of a gene, we chose the longest isoform of each gene, however in cases where isoforms had the same length, they were sorted alphabetically. This way, each m^6^A modification site has a unique gene ID, transcript (isoform) ID and functional category (5'UTR/CDS/3'UTR/intron or noncoding).

NGS data processing

Following the previous study, the sequencing reads were aligned to human genome (hg19) with STAR (32). The
default parameters were used. The read counts for each gene were counted using htseq-count (33) with default parameters. Only uniquely mapped reads were counted. The “expressed genes in HeLa cells” are defined as those genes with raw read count >50 in si-control cells.

**Differential splicing (DS) genes**

We used rMATS (34) to determine the DS genes in si-METTL3 or si-YTHDF1 vs. control. Note that mRNA-Seq rather than Ribo-Seq should be used for DS analysis. In the output file of rMATS software, if a splicing event has false discovery rate (FDR) <0.05 (35), it is recognized as a DS event, and the “host gene” of this DS event is termed DS gene. Accordingly, the “host exon” that contains the DS event is termed DS exon. There are five types of splicing according to the rMATS manual (Figure 1A). Any types of splicing with FDR <0.05 are regarded as DS events. When calculating the enrichment of DS events, since m\(^6\)A modifications are enriched in 3’UTRs and around stop codons, we should not directly count the number of DS events caused by si-METTL3. We took the m\(^6\)A distribution into consideration (as background). Take CDS as an example, we define (I) percentage of m\(^6\)A sites in CDS = number of m\(^6\)A sites in CDS/total number of m\(^6\)A sites on mRNA; (II) percentage of DS events in CDS = number of DS events in CDS/total number of DS events; (III) enrichment of DS events in CDS = percentage of DS events in CDS/percentage of m\(^6\)A sites in CDS.

**TE of genes**

We utilized TE = RPKM in Ribo-Seq/RPKM in mRNA-Seq (6,9,30) to determine the TE of each gene or the TE-foldchange between si-METTL3 or si-YTHDF1 vs. control libraries. RPKM stands for reads per kilobase per million (reads). Note that when calculating TE, only the reads mapped to CDS regions are counted.

**Statistical analyses**

We used R language to conduct statistical analyses (https://
www.R-project.org/). The comparisons of TE-foldchange values were performed using Wilcoxon rank sum tests. The enrichment comparison (between fractions) was performed using Fisher’s exact tests. The statistical significance was denoted as: *, P<0.05; **, P<0.01; ***, P<0.001.

Results

Splicing changes when METTL3 was knocked down were enriched around m\(^3\)A exons

There are five types of alternative splicing patterns, SE (skipped exon), A5SS (alternative 5’ splice site), A3SS (alternative 3’ splice site), MEX (multiple exclusive exon) and RI (retained intron) (Figure 1A). In the mRNA-Seq data of si-METTL3 vs. control, any types of splicing with FDR <0.05 are defined as DS events (Methods). The genes with DS events are defined as DS genes when METTL3 was knocked down.

Among the 24,035 expressed human genes in HeLa cells, 6,252 are m\(^3\)A genes and 17,783 are non-m\(^3\)A genes (Figure 1B). Interestingly, 486/6,252 (7.8%) out of the m\(^3\)A genes and 172/17,783 (0.97%) out of the non-m\(^3\)A genes are DS genes when METTL3 was knocked down (P value <0.001, Fisher’s exact test; Figure 1B). Furthermore, we found that 425/486 (87.4%) of the DS genes had m\(^3\)A modifications on the DS exon or the on the exons/introns flanking the DS exon (Figure 1B), indicating that the m\(^3\)A sites close to the DS exons might be responsible for the splicing changes.

DS events are relatively enriched in 5’UTRs

We questioned whether the DS events have preference in mRNA locations. We focus on the DS events in the 486 m\(^3\)A genes and calculated the relative enrichment of DS events in each mRNA location (Methods). We found that the enrichment of DS events in 5’UTR is greater than 1 while those of other categories (CDS, 3’UTR, intron or noncoding) do not significantly differ from 1 (Figure 2A). Our next question is based on the fact that many cis-elements in 5’UTR of mRNA play essential roles in determining the translation initiation, so could the DS events in 5’UTR cause the gain or loss of these essential cis-elements and consequently affect the translation of host genes?

DS events in 5’UTRs cause stronger alteration of TE

We profiled the TE foldchange of all genes in si-METTL3 vs. control. The TE of the 6,252 m\(^3\)A genes are globally down-regulated when METTL3 was knocked down (Figure 2B), which agrees with previous knowledge (6). Among the 486 DS m\(^3\)A genes, the genes with DS events in 5’UTR are highlighted against genes with DS events in other regions (CDS, 3’UTR and intron are combined) (Figure 2C). We could see that the gene with DS events in 5’UTR tend to have greater changes in TE (either increase or decrease). Furthermore, the absolute values of TE foldchange clearly show that the genes with DS events in 5’UTR alter their TE more severely than those with DS events in other regions (Figure 2D).

Translation of the 5’UTR DS genes is generally unaffected by YTHDF1

For the 425 m\(^3\)A genes with DS events near m\(^3\)A sites (Figures 1B, 6), 105 have DS events in 5’UTR (Figure 3). Our hypothesis is that the translation of these 5’UTR DS genes might be affected by additional cis elements, not only the reader protein alone (this does not mean the readers do not contribute). First, we should confirm that these 105 genes are not differentially spliced in YTHDF1-KD. 98 out of the 105 genes are non-DS genes in YTHDF1-KD (Figure 3), suggesting that the splicing changes observed in METTL3-KD are mainly caused by m\(^3\)A modifications rather than reader proteins.

We next investigated the TE foldchange in YTHDF1-KD vs. control. Different gene sets showed distinct patterns (Figure 3). For the 17,783 non-m\(^3\)A genes (background), their TE log₂-foldchanges had a median value around zero, which agreed with our expectation. For the 5,766 non-DS m\(^3\)A genes (in METTL3-KD), their TEs were significantly down-regulated in YTHDF1-KD (Figure 3). The same trend of decreased TE went for the 320 DS m\(^3\)A genes which had DS events in non-5’UTR regions (Figure 3). This indicated that m\(^3\)A genes would be translationally down-regulated when reader protein is knocked down. For the 98 DS genes in 5’UTR, their TE was only slightly and insignificantly down-regulated in YTHDF1-KD (Figure 3). This result proves that the TE changes of the 98 [103] genes observed in METTL3-KD were majorly contributed by DS-related cis changes. The reader protein alone could not account for such a big change (Figure 2C,D) as demonstrated in Figure 3.

Discussion

RNA modifications like the m\(^3\)A methylation is highly
regulated by different factors and meanwhile regulating its downstream biological processes. As we have introduced, the dynamic regulation of m^6A modification is accomplished by the writers (13,14) and erasers (15,16), and the downstream effects of m^6A are exerted with the help of readers (5,6,11,17,36) or by m^6A site itself (1,4). However, it is not strange at all that the effects of “readers” and “m^6A site itself” could simultaneously contribute to the translation status of host genes. Our goal is to search for the minor and indirect effects of m^6A site itself on the translation of host genes.

By comparing the global changes in mRNA splicing patterns and TE of si-METTL3 vs. control in human HeLa cells, we found that the DS genes are enriched in m^6A modified genes, suggesting that the DS events (in si-METTL3 vs. control) are likely caused by m^6A modifications. Changes in splicing patterns are often related to 5’UTR. The global TE of m^6A modified genes (no matter the modification is on UTR or CDS) is down-regulated when METTL3 was knocked down. This result agrees with previous knowledge that m^6A enhances the translation of host gene by the reader protein (6). We further found that the genes with DS events in 5’UTR alter their TE more severely than those genes with DS events in CDS/3’UTR/intron, suggesting a role of m^6A-mediated splicing changes in translation regulation. Importantly, we verified our hypothesis in the YTHDF1-KD samples. The TE of the few 5’UTR DS genes in METTL3-KD was unchanged in YTHDF1-KD (Figure 3), indicating that their TE changes in METTL3-KD might be largely due to the effect of DS caused by m^6A itself.

The established theory of how m^6A affects mRNA translation includes (I) YTHDF1 recruits the initiation factors to m^6A modified mRNAs and promote the translation (6) and (II) m^6A on 5’UTR acts as IRES to

Figure 2 The TE foldchange of genes in si-METTL3 vs. control. (A) Enrichment of DS events compared to the distribution of m^6A sites in DS genes. Fisher’s exact test; (B) TE of m^6A (red) and non-m^6A genes; (C) dot plot displaying TE of m^6A genes with DS events in 5’UTR (orange) and other regions; (D) barplot displaying TE foldchange of m^6A genes with DS events in 5’UTR and other regions. The absolute value of log_2 TE-foldchange is shown. Error bar represents stand error. Wilcoxon rank sum test. **, P<0.01. m^6A, N^6-methyladenosine; TE, translation efficiency; DS, differential splicing.
directly initiate translation (9,30). In contrast to these big findings by previous literatures, our analysis suggests the fine-tuning of mRNA translation mediated by \( m^6A \) modification and splicing changes. These fine-tuning effects are indirect and relatively weaker than those direct approaches.

One might think that “independent of reader proteins” could be a new mechanism. However, we do not wish to overstate this conclusion. It only account for a few genes that have 5’UTR \( m^6A \) sites and meanwhile have DS events in 5’UTR when \( m^6A \) is knocked down (conservatively 98 genes). Let us take the “microRNA binding to 3’UTR” as an example. Presume that an \( m^6A \) site is located in 3’UTR of a gene and the suppression of methylation results in DS in 3’UTR region. If this DS event in 3’UTR leads to the gain or loss of microRNA binding sites, then it might eventually cause changes in translation or mRNA expression of host gene. We think this example could not be called a new mechanism because (I) “\( m^6A \) affects splicing” is known; (II) “Splicing might cause gain or loss of microRNA binding site” is conceivable; (III) “microRNA binding affects host gene translation or expression” is known. There are only a few candidate genes that could fit all these criteria. This pattern is merely valid for specific genes rather than all genes (just because these few gene happened to fit the criteria). Similarly, our “\( m^6A\)-splicing-5’UTR-translation” chain is based on known mechanisms in every step. Only a small set of genes could link all these steps together (conservatively 98 genes in Figure 3). Our purpose is to reveal this observation and show that the \( m^6A \) modification could affect the host gene through this indirect way.

Next, we want to clarify that our results do not conflict with known theories. The translational effect caused by 5’UTR DS events is “independent of reader proteins” does not mean that the reader proteins contribute nothing. We should clearly clarify that “both the 5’UTR DS events and the reader protein contribute to the observed translational changes”. The YTHDF1-dependent pathway to enhance
the translation of m^6A genes was discovered years ago (6). We think that the translational changes are caused by multiple factors. It is like the multiple regression analysis. The global trend showed that the TE of m^6A genes were down-regulated in YTHDF1-KD (6), which proved the positive contribution of YTHDF1. For some m^6A genes with up-regulated TE, the elevated TE might be explained by other variables (like the gain or loss of some cis-elements in UTRs). Similarly, in our analyses, we found 98 genes that their DS events strongly affects their translation and the reader protein YTHDF1 plays a less important role. This does not mean that YTHDF1 does not contribute to the translation of target genes. Thus, our findings do not conflict with known theories.

Our work might remind people that the translation status is affected by numerous factors and none of the factors alone could explain all the observed translational changes. Although the global comparison showed that thousands of m^6A genes were translationally down-regulated when YTHDF1 was knocked down (6), one could always find the exceptions that a few m^6A genes were up-regulated, which might be caused by other factors including cis elements in UTRs. At this stage, the detailed mechanism of how DS events affect the translation of host gene remains unexplored. What we know is that there are many cis-elements in 5'UTR that determine the translation initiation, and that the DS events in 5'UTR might cause the gain or loss of these important cis-elements and eventually impact the translation of downstream CDS.

**Conclusions**

Our results reveal that m^6A modifications could modulate the translation of mRNAs through affecting their splicing patterns. The differential splicing caused by m^6A are enriched in 5'UTR and consequently affect translation. These indirect effects are independent of the direct regulation by reader proteins, and likely to be related to the cis-elements in 5'UTR that determine the translation of downstream CDS. Our work broadened our knowledge about the translation regulation by m^6A modifications.

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**Footnote**

**Conflicts of Interest:** All authors have completed the ICMJE uniform disclosure form (available at http://dx.doi.org/10.21037/tcr.2019.09.18). The authors have no conflicts of interest to declare.

**Ethical Statement:** The authors are accountable for all aspects of the work in ensuring that questions related to the accuracy or integrity of any part of the work are appropriately investigated and resolved. This study was conducted in accordance with the Declaration of Helsinki (as revised in 2013). Institutional ethical approval and individual informed consent were waived.

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