RNA-binding proteins related to stress response and differentiation in protozoa

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Abstract

RNA-binding proteins (RBPs) are key regulators of gene expression. There are several distinct families of RBPs and they are involved in the cellular response to environmental changes, cell differentiation and cell death. The RBPs can differentially combine with RNA molecules and form ribonucleoprotein (RNP) complexes, defining the function and fate of RNA molecules in the cell. RBPs display diverse domains that allow them to be categorized into distinct families. They play important roles in the cellular response to physiological stress, in cell differentiation, and, it is believed, in the cellular localization of certain mRNAs. In several protozoa, a physiological stress (nutritional, temperature or pH) triggers differentiation to a distinct developmental stage. Most of the RBPs characterized in protozoa arise from trypanosomatids. In these protozoa gene expression regulation is mostly post-transcriptional, which suggests that some RBPs might display regulatory functions distinct from those described for other eukaryotes. mRNA stability can be altered as a response to stress. Transcripts are sequestered to RNA granules that ultimately modulate their availability to the translation machinery, storage or degradation, depending on the associated proteins. These aggregates of mRNPs containing mRNAs that are not being translated coloc- alize in cytoplasmic foci, and their numbers and size vary according to cell conditions such as oxidative stress, nutritional status and treatment with drugs that inhibit translation.

Key words: Gene expression regulation; RNA-binding proteins; RNA-protein complexes; RNA granules; Protozoa; Stress and cell differentiation

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Core tip: RNA-binding proteins (RBPs) are numerous and widely distributed in nature. In addition to having different domains, these proteins are key modulators of gene expression and are involved in the cellular response to environmental changes, cell differentiation and cell death. In protozoa RBPs are crucial for the rapid gene expression remodeling that occurs in the course of cell differentiation or the stress response.

INTRODUCTION

RNA-binding proteins (RBPs) are numerous and widely distributed in nature and are involved in gene expression control at all levels. There are several distinct families of RBPs. They are key modulators of gene expression and are involved in the cellular response to environmental changes, cell differentiation and cell death. The RBPs can differentially combine with RNA molecules and form ribonucleoprotein (RNP) complexes, thereby defining the function and fate of the RNA molecules within the cell.

RBPs and the RNA-binding domains

RBPs have multiple and diverse domains; many of them have a combinatorial set of RNA-binding domains (RBDs) that allow various modes of recognition to exert their function in the cell. The RBD recognizes and interacts with the RNA molecule based on sequence and/or structure. In addition to the RBD domain, many of the RBPs also have protein-protein domains that form complexes that are also subjected to intense post-translational modifications, such as methylation and phosphorylation, further increasing the complexity of the study and characterization of RBPs.

One of the most common domains found in RBPs is the RNA-recognition motif (RRM); it shows affinity with single-stranded RNA, DNA and proteins. Approximately 90 amino acid residues are involved in the recognition and binding of a target RNA sequence of 2 to 6 nucleotides. The proteins that contain this domain are involved in all pathways of the RNA cycle, from splicing to mRNA turnover. The RRM is usually present in multiple copies within a protein, and this multiplicity enhances the specificity of the ligand.

The zinc finger domain was first described as a DNA-binding domain (with the CCHH amino acid motif) and was subsequently described in proteins that interact with RNA (with the CCCH motif), such as Tis11 and Tristetraprolin (TTP). TTP is involved in transcript degradation through its interaction with the AU-rich elements in the 3′-untranslated regions (3′-UTRs) of some mRNAs, as described for tumor necrosis factor (TNF)-responsive factor alpha and granulocyte-macrophage colony-stimulating factor mRNAs.

The Pumilio (PUF) protein takes part in regulating different processes in the cell, such as embryogenesis, development, and differentiation; however, the best-characterized function of PUF is as a post-transcriptional repressor. The domain is characterized by a highly conserved C-terminal RBD composed of eight tandem repeats.

The hnRNP K-homology domain recognizes a specific sequence of single-stranded RNA and is found in both eukaryotes and prokaryotes. The domain is approximately 70 amino acids long, and the proteins are associated with a diversity of gene expression regulation mechanisms, such as splicing, transcriptional regulation, and translational control.

The Arg-Gly-Gly repeat (RGG) domain can interact with either RNAs or proteins, and this interaction can be based on the positive charge of the arginine amino acids. Proteins containing RGG motifs are associated with several cellular processes, including transcription, splicing and mRNA export. One example of an RGG domain protein is the yeast Scd6 protein. It modulates translation by preventing the formation of the 48S pre-initiation complex by binding eIF4G through its RGG motif.

RBPs in protozoa

Most of the RBPs characterized in protozoa arise from trypanosomatids. Proteins such as TcDHH1 are involved in the formation of mRNA granules in Trypanosoma cruzi and Trypanosoma brucei, but it is believed that the function and composition of these granules can be different from those described for higher eukaryotes. TcDHH1 binds to RNAs encoding stage-specific surface antigens from the infective trypanomastigote forms. However, these mRNAs are also found at low levels in the non-infective epimastigote forms, leading to the hypothesis that TcDHH1 might be involved in the degradation of the targets in these forms.

Trypanosome gene expression regulation is mostly post-transcriptional, which suggests that some RBPs might display regulatory functions distinct from those described for other eukaryotes. One example is the poly-A binding proteins PABP1 and PABP2: Only PABP2 accumulates in the nucleus, indicating that PABP1 and PABP2 might be part of distinct mRNPs that associate with distinct populations of mRNAs. Some proteins, such as TcDHH3, TcPTB2, TcZPF3, TcPUF9, TcPUF1 and TcPUP2, can function as mRNA destabilization factors. However, the modulation mechanisms for most RBPs in trypanosomatids remain unknown.

In T. cruzi, the zinc finger protein TcZFP1 binds preferentially to C-rich regions. However, TcZFP2 has a higher affinity for A-rich regions, and the mRNAs associated with this protein appear to be upregulated in metacyclic trypomastigotes, suggesting a possible role in parasite differentiation.

Some RBPs, such as TcPuf6 and TcUbp1, are involved in the downregulation of their target transcripts, while others, such as RBP19, can negatively regulate their own mRNAs. Some RBPs, such as RBP42 and the Alba proteins, associate with the translation machinery. The RBP42 protein from T. brucei binds to the coding region of mRNAs of the insect form that encode metabolic proteins. RBP42 is an essential protein and associates with mRNAs bound to polysomes in the cytoplasm. However, some RBPs, such as...
TbRBP10, a cytoplasmic protein essential to the *T. brucei* bloodstream forms, appear to affect the expression of their targets indirectly through other proteins that form the mRNP complex\(^{22}\).

TcRBP40 is found dispersed within the cytoplasm and is also concentrated in reservosomes, a storage organelle present in *T. cruzi* and some bat trypanosomes during a specific life cycle stage\(^{23,24}\), suggesting that this organelle can also be involved in RNA processing\(^{25}\). There are also some RBPs with assigned nuclear functions, such as TbPTB2, which is involved in translancing processing of some mRNAs\(^{26}\). Similarly, TbRRM1 is a nuclear RBP that associates with mRNAs from *T. brucei* and with the auxiliary splicing factor polypyrimidine tract-binding protein (PTB) 2, but not with components of the core spliceosome. A knockdown of TbRRM1 caused extensive alterations in mRNA abundance and regions enriched for the downregulated mRNAs were identified. In addition, in cells subjected to heat shock, TbRRM1 shifted from the nucleus to the cytoplasm, which led to the compaction of chromatin\(^{27}\).

The *T. brucei* nucleolar RBPs PUF7 and PUF10 are involved in ribosomal RNA maturation and transcriptional control of procyclic stage-specific genes transcribed by RNA polymerase I \(^{28}\). TcSUB2 is involved in the transcription/export pathway and is mainly localized to the nucleus\(^{29}\).

In *Plasmodium*, DDX-6 class DEAD box RNA helicase (DOZI) is essential to zygote development because of its role in controlling mRNA stability\(^{30,31}\). In *P. falciparum*, both PfPuf1 (PF003935c) and PfPuf2 (PF00825c) are upregulated in gametocytes\(^{32,33}\). Microarray analysis detected increasing amounts of PfPuf1 and PfPuf2 mRNAs during gametocyte maturation\(^{34}\). Interestingly, the highest level of PfPuf2 expression was observed in sporozoites\(^{35}\).

In *Toxoplasma gondii*, TgPuf1 displays different expression levels in tachyzoites and bradyzoites, indicating that it might function in regulating processes such as proliferation and/or differentiation that would enable the parasites to respond rapidly to changes in environmental conditions\(^{36}\). Another protein from *T. gondii* is the RGG single-strand-binding protein (TgSsossB), which interacts with the TgAlba proteins that are involved in translation regulation\(^{37}\). When the RGG domain was deleted from the protein, the mutant strain produced fewer plaques in stress conditions, a defect associated to a slow growth phenotype due to the exposure of extracellular parasites to stress. Moreover, the mutant lost its capacity to interact with the TgAlba complex\(^{38}\).

**RBPs and post-transcriptional control in stress**

In unicellular and multicellular organisms, there is a very subtle balance between the cell’s capacity to respond to stress and cell death. For the former, this balance needs to be even more strictly regulated, and there are many variables involved that take into account the strength and duration of the stimulus\(^{39}\). The stress response needs to be fast to allow cell survival, if we consider the cell’s time scale, at the moment the stress is triggered, the response starts immediately (within nanoseconds), and the cell metabolism adapts dynamically to the environment. Accordingly, there is modulation of gene expression, and trans-acting factors such as RBPs are key to this prompt response.

Post-transcriptional gene regulation occurs at different levels: Splicing, mRNA stability and translation regulation. With regard to splicing, it has been shown that when a cell is subjected to a low dose of ultraviolet radiation irradiation, the C-terminal domain of RNA polymerase II is phosphorylated, and the splicing dynamics are altered. This alteration allows, via alternative splicing, the inclusion of exons with weaker splice sites\(^{40,41}\). One example of alternative splicing occurs with the MDM2 RNA during genotoxic stress. Under normal conditions, the MDM2 protein targets the p53 protein for degradation via ubiquitination; however, during stress, an exon is skipped in the MDM2 RNA, which ultimately results in the maintenance of p53 in the cell. This effect is only reversed upon restoration of the normal condition\(^{42}\). Splicing factors are also targets of regulation during stress. For example, hnRNPI shuttles to the cytoplasm during osmotic stress, while the other splicing factors remain in the nucleus\(^{43}\).

Extensive studies have been performed and highlight the importance of translational control during stress. For the cell, it is vital to control and arrest translation during stress, as 50% of the cell’s energy is consumed during translation\(^{44}\).

For most mRNAs, translation initiation occurs when the trimeric eIF4F complex binds to the transcript. This complex is formed by the protein eIF4E (the cap-binding protein), eIF4A (RNA helicase) and eIF4G (which forms the scaffold protein complex via its binding sites for eIF4E and eIF4A). The trimeric complex binds to the cap, followed by ribosomal scanning from the RNA 5'-end to the first AUG site. Then, eukaryotic initiation factor 2 (eIF2) associates with tRNAmet, part of eIF2-GTP ternary complex that is required to bring the initiator tRNA to the ribosome\(^{45,46}\). When a stress signal is generated, two basic mechanisms are activated: The availability of eIF4E, which can be sequestered by eIF4EBPs (eIF4E-binding proteins), and the phosphorylation of eIF2a. Global translational arrest occurs, and only a few proteins are translated to help the cell overcome the stress. The most studied genes upregulated during stress are those encoding heat shock proteins (HSPs), whose transcription is stimulated by the heat shock factor that senses misfolded proteins in the cytoplasm and shuttles to the nucleus. HSPs are chaperones that help properly fold proteins\(^{47}\). Aside from HSPs, some transcripts that contain upstream open reading frames can be translated even under conditions of eIF2a phosphorylation\(^{48,49}\). The transcripts that contain internal ribosome entry sites (IRESs) on their 5'-UTRs can also bypass the inhibition of cap-dependent control (eIF4F). Interestingly, some
formation of these granules. Some of the proteins characteristic of P-bodies are depicted. There is a spatial separation between the mRNAs that are being translated from those that are associated with P-bodies. The mRNA translation status depends on the proteins associated with the mRNP complexes, which direct the mRNA molecules either to the polysomes for translation or to RNA granules for storage and/or degradation. Observations indicate that several mRNAs present in the P-bodies undergo cap removal and are consequently prone to degradation. However, the P-bodies may also act as mRNA storage sites, allowing mRNAs to return to the translation machinery. Experiments have shown that specific mRNAs accumulate in P-bodies in growing cells or cells under stress and that they emerge from the granules and migrate to the polysomes to be translated when normal conditions are restored. Another type of mRNP granule found in eukaryotes is the stress granule. Unlike P-bodies, the stress granules are composed mainly of translation initiation factors, the 40s ribosomal subunit and several RBPs. Studies indicate that physical interaction occurs between the P-bodies and stress granules and that the mRNAs can switch between them (Figure 1). The translation initiation complexes bound to mRNAs, which include eIF3, the eIF4F complex, eIF4B, the 40S ribosomal subunit and PABP-1, form the core of the stress granules and are considered markers of these...
structures. Other proteins characteristic of stress granules may affect mRNA translation, such as T cell antigen internal-1 (TIA-1) and TIA-1-related, Fragile X mental retardation protein (FMRP) and Fragile X mental retardation-related protein-1, Argonaute, cytoplasmic polyadenylation element-binding protein, Pumilio and RNA-associated protein 55. Still others are involved in mRNA stability, such as RNA helicase RCK/p54, 5'-exonuclease 3' XRN1, proteins that bind to the ARE element, and Hur and TTP, which respectively stabilize and destabilize mRNA. In mammalian cells, stress granules are formed as a result of phosphorylation of eIF2α by stress-activated kinases. The granules assemble when RBPs such as TIA-1, TTP, FMRP or G3BP bind to specific transcripts that are linked to the stalled pre-initiation 48S complex. After this initial nucleation stage, the polyA-binding protein (PABP) promotes the aggregation of these small granules by making them microscopically visible. Some proteins that do not have a canonical RBP domain can also associate with the granules through protein-protein interactions, which suggests that stress granules can interact with other signaling pathways in the cell.

Sheth et al. proposed an RNA cycle model describing the dynamics of mRNAs in the cytoplasm, from polysomes to P-bodies and stress granules. In this model, mRNAs are able to associate with polysomes being effectively translated into proteins; however, when a stress signal is triggered, mRNAs interact with components of the degradation/storage pathway, translation is stopped, and the mRNP that is formed is held in P-bodies. This mRNA can then be stored or degraded in the P-bodies or returned to the translation apparatus. Stress granules represent a translation arrest site that increases in abundance as mRNAs are dissociated from polysomes; the association of mRNA with initiation factors may facilitate their re-entry for translation when the stress ceases. The molecular mechanisms encompassed in the transition between polysomes, P-bodies and stress granules need to be clearly elucidated.

In several unicellular eukaryotes, stress is involved in triggering differentiation. This association is exemplified in Figure 2, which shows that nutritional, pH and temperature stresses are involved in the life cycle of the human pathogen Trypanosoma cruzi. Accordingly, it has been shown that formation of RNA granules increases during the nutritional stress preceding metacyclogenesis. Stress and differentiation

Figure 2 Physiological stresses and differentiation. The schematic representation shows the life cycle of Trypanosoma cruzi. This parasite alternates between two distinct intermediary hosts. The change to each host involves a temperature stress, either a heat shock (28 °C to 37 °C) during the passage from the invertebrate to the vertebrate or a cold shock (37 °C to 28 °C) when the parasite switches from the mammalian to the triatomine vector. Inside the hosts, nutritional and pH stresses are determinants for the differentiation from the non-replicative to the replicative forms (either epimastigotes in the insect or amastigotes in the mammal host) and from those to the infective trypomastigote forms.

**RBP s in stress in protozoa**

RBPs are crucial for the rapid gene expression remodeling that occurs in the course of cell differentiation or...
the stress response. In some case, the changes in gene expression are accompanied by changes in mRNA localization within the cell. Accordingly, the localization of specific mRNAs is a meaningful tool to optimize protein synthesis and to maintain cell polarity\(^{[69]}\). Fluorescence in situ hybridization analysis using poly-T showed the increase of fluorescent granules during stress conditions in trypanosomatids, suggesting that the mRNAs were mobilized to form RNA granules. It was also observed that the cytoplasmic localization of specific transcripts changed during differentiation\(^{[70]}\).

Proteomic analysis evidenced a global change in the protein composition of \textit{T. brucei} mRNPs isolated from polysomal and post-polysomal fractions from epimastigotes in exponential growth and epimastigotes under nutritional stress. More than 500 proteins were identified, and among them were several RBPs. Interestingly, a dynamic shift in protein composition was also observed in response to nutritional stress, especially for those proteins identified in the polysomal-free fraction of stressed parasites\(^{[71]}\).

Some specific RBPs play important roles during the stress response in parasitids. One example is TcZC3H39, a zinc finger protein from \textit{T. cruzi} that exhibits a shift in the composition of the bound mRNA targets depending on the physiological conditions of the cell. Under stress conditions, the TcZC3H39-mRNP complex sequesters highly expressed mRNAs, thereby slowing translation activity. Accordingly, the number of bound mRNA targets was higher in stressed parasites than in non-stressed parasites. Under stress conditions, the TcZC3H39-mRNP complex was enriched in targets that are highly expressed under non-stress conditions, such as ribosomal and oxidative phosphorylation pathway proteins\(^{[68]}\). The protein content of TcZC3H39-mRNP consists of ribosomes, translation factors, RNA helicases and other RBPs, suggesting that this zinc finger protein might be part of complexes similar to stress granules, sequestering highly expressed mRNAs and the associated ribosomes, possibly slowing down translation in response to stress conditions\(^{[68]}\).

The zinc finger protein ZC3H11 plays an important role during \textit{T. brucei} heat shock. This protein is essential for the bloodstream form of the parasite and stabilizes HSP70 mRNA and other transcripts that encode chaperones necessary for proper protein folding or refolding during heat stress\(^{[72]}\). ZC3H11 interacts with the Mkt1 and PBP1 proteins, forming a complex that stabilizes the associated transcripts\(^{[73]}\).

TbDRBD3 is a cytoplasmic RRM protein from \textit{T. brucei} that relocates to the nucleus during actinomycin-D treatment\(^{[76]}\). Other RBPs, such as DRBD4 (PTB2 in \textit{T. brucei}, a PTB homolog) and PABP, have also been observed to show this behavior during treatment with actinomycin-D or during severe heat shock\(^{[76]}\). It has been hypothesized that the nucleolus plays a role as a sensor or regulator of cell metabolism during the stress response\(^{[77,78]}\).

\textit{T. brucei} Alba 3 and 4 are found in stress granule-like structures during nutritional deprivation. They partially co-migrate with polysomes and also associate with the helicase DHH1, translation initiation factors and PABP1, indicating a role in translation modulation\(^{[21]}\).

The heat shock response in \textit{T. brucei} involves a reduction of the polysomes and a corresponding increase in the number of RNA granules, such as P-bodies and stress granules. The P-bodies from \textit{T. brucei} are composed of XRNA (mRNA decay machinery), Scd6 and DHH1. These RNA granules could act as sites for RNA storage or degradation during stress and differentiation\(^{[79]}\).

In \textit{T. cruzi}, the DHH1 protein is also localized in cytoplasmic foci that resemble P-bodies and/or stress granules. The protein interacts with PABP, and its localization changes in response to nutritional stress; the granules increase in number and also become bigger, likely due to the shift of mRNAs from the polysomes to the RNA granules\(^{[12]}\).

It is important to mention that proteins lacking canonical RBDS but having other functions in the cell can also associate with RNAs and become structural components of RNP complexes\(^{[68]}\). The moonlighting theory proposes that some multifunctional enzymes normally associated with metabolic pathways can also play a role in regulating gene expression in prokaryotes and eukaryotes. This “moonlighting” role was previously described for glyceraldehyde-3-phosphate dehydrogenase, which can be found within the nuclei of mammalian cells acting in transcription, DNA repair, RNA binding and transport, and telomere binding\(^{[73,81,82]}\). Other examples are inosine monophosphate dehydrogenase\(^{[83]}\), acetyl-CoA carboxylase\(^{[84]}\), lipin-1\(^{[85]}\), aminocyl-tRNA synthetase\(^{[86]}\), Lv5p, which is required for the synthesis of branched chain amino acids\(^{[87]}\), phosphofructokinase\(^{[88]}\) and the mitochondrial enzyme Arg5,6, that participates in the transcription of target genes in addition to its role in arginine biosynthesis\(^{[89]}\).

Another example of a moonlighting protein in \textit{T. cruzi} is EF-1\(\alpha\), which associates with a specific subset of mRNAs during stress conditions. This protein can associate with large protein complexes independent of the translation machinery. A specific subset of mRNAs associates with EF-1\(\alpha\)-mRNPs in unstressed or stressed epimastigotes. Some mRNAs are common to both physiological conditions, whereas others are specific to a given physiological condition. Gene ontology analysis identified enrichment of gene sets involved in single-organism metabolic processes, amino acid metabolic processes, ATP and metal ion binding, glycolysis, glutamine metabolic processes, and cobalt and iron ion
binding\textsuperscript{(39,40)}.  

RBPs and eukaryotic cell differentiation  

In parasites, several RBPs involved in differentiation have been described. The RRM-containing domain proteins RBP10 and RBP6 from T. brucei are implicated in life-cycle differentiation\textsuperscript{(22,61)}. Accordingly, RBP10 knockdown led to a great alteration in the mRNA levels of transcripts normally found in high levels in the bloodstream forms\textsuperscript{(22)}. The CCCH zinc finger proteins ZFP1, ZFP2 and ZFP3 have also been described as having roles in differentiation in trypanosomes\textsuperscript{(22,59,59)}. Knockdown of ZFP1 disturbed the bloodstream differentiation by repositioning the kinetoplast in T. brucei\textsuperscript{(60)}. A similar result was obtained with the RNAi knockdown for ZFP2: The parasite was unable to differentiate\textsuperscript{(92,93)}, ZFP3 overexpression enhanced the differentiation levels to procyclic forms, and it is believed that these three proteins are part of the same differentiation pathway\textsuperscript{(93,65)}.  

CONCLUSION  

Despite evidence of the role for RBPs in gene expression regulation, particularly in protozoa as a means to respond promptly to environmental changes, the detailed mechanisms remain to be clearly elucidated. We anticipate that the complexity of these mechanisms should increase, as we do not yet know how different combinations of RBPs in a given complex affect the fate of the bound mRNAs. Furthermore, the modulation of the interactions within a complex could also be affected by post-translational modifications of the RBPs, creating an RBP code analogous to the histone code.  

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