

The exon junction complex is detected on CBP80-bound but not eIF4E-bound mRNA in mammalian cells: dynamics of mRNP remodeling

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Newly spliced mRNAs in mammalian cells are characterized by a complex of proteins at exon–exon junctions. This complex recruits Upf3 and Upf2, which function in nonsense-mediated mRNA decay (NMD). Both Upf proteins are detected on mRNA bound by the major nuclear cap-binding proteins CBP80/CBP20 but not mRNA bound by the major cytoplasmic cap-binding protein eIF4E. These and other data indicate that NMD targets CBP80-bound mRNA during a ‘pioneer’ round of translation, but whether nuclear eIF4E also binds nascent but dead-end transcripts is unclear. Here we provide evidence that nuclear CBP80 but not nuclear eIF4E is readily detected in association with intron-containing RNA and the C-terminal domain of RNA polymerase II. Consistent with this evidence, we demonstrate that RNPS1, Y14, SRm160, REF/Aly, TAP, Upf3X and Upf2 are detected in the nuclear fraction on CBP80-bound but not eIF4E-bound mRNA. Each of these proteins is also detected on CBP80-bound mRNA in the cytoplasmic fraction, indicating a presence on mRNA after export. The dynamics of mRNP composition before and after mRNA export are discussed.

Keywords: cap-binding proteins/CTD/exon junction complex/mRNP transport/Upf proteins

Introduction

In mammalian cells, the expression of protein-encoding genes requires a series of steps in which pre-mRNA is processed to mRNA in the nucleus before mRNA is translated into protein in the cytoplasm. These steps are subject to quality control to ensure that only completely processed mRNA is exported to the cytoplasm (reviewed in Maquat and Carmichael, 2001). An additional quality control, called mRNA surveillance or nonsense-mediated mRNA decay (NMD), degrades mRNAs that prematurely terminate translation >50–55 nucleotides upstream of an exon–exon junction as a means to prevent the synthesis of potentially harmful truncated proteins (reviewed in Maquat, 1995, 2000, 2002; Li and Wilkinson, 1998; Culbertson, 1999; Hentze and Kulozik, 1999; Wilusz and

Peltz, 2001). Beside translation, NMD in mammalian cells requires at least four proteins: Upf1, Upf2, Upf3/3X (related products of different genes that are also called Upf3a/b) and hSMG1/ATX (Sun *et al.*, 1998; Lykke-Andersen *et al.*, 2000; Mendell *et al.*, 2000; Denning *et al.*, 2001; Serin *et al.*, 2001; Yamashita *et al.*, 2001; K.M.Brumbaugh, D.M.Otterness, X.Li, F.Lejeune, R.S.Tibbetts, L.E.Maquat and R.T.Abraham, unpublished data). Studies of Upf orthologs in *Saccharomyces cerevisiae* indicate that Upf1 interacts with eukaryotic release factors (eRFs) 1 and 3, Upf2 and Upf3 interact with eRF3 in a way that competes with the eRF3–eRF1 interaction, and all three proteins influence translation termination efficiency (Czapinski *et al.*, 1998; Maderazo *et al.*, 2000; Wang *et al.*, 2001). hSMG1/ATX, like its ortholog in *Caenorhabditis elegans* (Page *et al.*, 1999), is a phosphatidylinositol 3-kinase-related protein kinase involved in the phosphorylation of Upf1 (Denning *et al.*, 2001; Pal *et al.*, 2001; Yamashita *et al.*, 2001; K.M.Brumbaugh, D.M.Otterness, X.Li, F.Lejeune, R.S.Tibbetts, L.E.Maquat and R.T.Abraham, unpublished data).

Exon–exon junctions have been proposed to function in NMD via the ~335 kDa exon junction complex (EJC) of proteins that is deposited ~20–24 nucleotides upstream of junctions as a consequence of pre-mRNA splicing (Le Hir *et al.*, 2000a,b, 2001a; Kataoka *et al.*, 2001; Kim *et al.*, 2001b; Lykke-Andersen *et al.*, 2001). Components of this complex include REF/Aly, Y14, DEK, SRm160 and RNPS1. REF/Aly facilitates the nuclear export of mRNA by interacting with the mRNA export receptor TAP (Katahira *et al.*, 1999; Lou and Reed, 1999; Bachi *et al.*, 2000; Kataoka *et al.*, 2000, 2001; Stutz *et al.*, 2000; Zhou *et al.*, 2000; Le Hir *et al.*, 2001a; Rodrigues *et al.*, 2001). Y14, which binds to mRNA that has undergone splicing (Kataoka *et al.*, 2000) and interacts with REF/Aly and RNPS1 *in vitro* (Kataoka *et al.*, 2001), has been proposed to provide a position-specific memory of the EJC in the cytoplasm since it is detected in association with both nuclear and newly exported cytoplasmic mRNA (Kim *et al.*, 2001b). DEK has multiple functions that include interacting with SR proteins during splicing (McGarvey *et al.*, 2000) as well as altering the superhelical density of DNA in chromatin (Alexiadis *et al.*, 2000) and altering the transcription of certain genes (Faulkner *et al.*, 2001). SRm160 co-activates pre-mRNA splicing (Blencowe *et al.*, 1998; Kataoka *et al.*, 2000; McGarvey *et al.*, 2000) and promotes transcript 3'-end cleavage (McCracken *et al.*, 2002). Notably, neither DEK nor SRm160 shuttle to the cytoplasm in assays using mammalian cell heterokaryons (Lykke-Andersen *et al.*, 2001; Y.Ishigaki, B.Blencowe and L.E.Maquat, unpublished data). RNPS1 functions in pre-mRNA splicing (Mayeda *et al.*, 1999) and recently was shown to connect splicing and NMD mechanistically

(Lykke-Andersen *et al.*, 2001): (i) RNPS1 and, to a lesser extent, Y14 tethered to the 3'-untranslated region of β -globin mRNA recapitulates the function of the EJC in NMD as does tethered Upf1, Upf2 or Upf3/3X (Lykke-Andersen *et al.*, 2000, 2001); and (ii) FLAG-RNPS1 transiently expressed in HEK293 cells co-immunoprecipitates with Upf1, Upf2 and Upf3/3X (Lykke-Andersen *et al.*, 2001). Considering that Upf3/3X, RNPS1 and Y14 are mostly nuclear but shuttle, Upf2 is cytoplasmic but primarily perinuclear, and Upf1 is primarily cytoplasmic (Lykke-Andersen *et al.*, 2000, 2001; Serin *et al.*, 2001; J.T.Mendell and H.C.Dietz, personal communication), these data indicate that Upf3/3X joins the splicing-dependent mRNP complex in the nucleus by interacting either directly or indirectly with RNPS1 and, possibly, Y14 (Lykke-Andersen *et al.*, 2001). Extending the idea that Upf3/3X is recruited by the complex, Y14 has been shown to interact with REF/Aly, TAP and Upf3/3X independently of RNA, and Upf3X has been shown to map *in vivo* upstream of the exon-exon junction of two spliced mRNAs (Kim *et al.*, 2001a). According to current thinking, Upf2 joins the complex during or immediately after export to the cytoplasm. Provided that translation terminates prematurely (i.e. >50–55 nucleotides upstream of an EJC-marked exon-exon junction), Upf1 subsequently interacts with the complex in a way that elicits NMD (Ishigaki *et al.*, 2001; Lykke-Andersen *et al.*, 2001).

Another important connection between splicing and NMD was elucidated recently with the finding that Upf2 and Upf3X can be detected on spliced mRNP immunopurified using anti-CBP80 antibody but not anti-eIF4E antibody (Ishigaki *et al.*, 2001). This result implies that the two Upf proteins are recruited to newly synthesized mRNA that has yet to be translated. CBP80 is the mostly nuclear cap-binding protein that shuttles in association with mRNA to the cytoplasm (Visa *et al.*, 1996; Shen *et al.*, 2000; Ishigaki *et al.*, 2001), where it is replaced by the mostly cytoplasmic cap-binding protein eukaryotic initiation factor (eIF) 4E (reviewed in Gingras *et al.*, 1999). CBP80-bound mRNA, rather than eIF4E-bound mRNA, was found to be the primary template for the first so-called 'pioneer' round of translation since it is the primary substrate of NMD (Ishigaki *et al.*, 2001). The conclusion that CBP80-bound mRNA is targeted for NMD was based on several criteria: (i) nonsense-containing CBP80-bound mRNA is reduced in abundance to an extent that is comparable to nonsense-containing eIF4E-bound mRNA, and CBP80-bound mRNA is thought to be a precursor to eIF4E-bound mRNA; (ii) the nonsense-mediated reduction in the abundance of CBP80-bound mRNA is abrogated by either a suppressor tRNA that recognizes the nonsense codon as encoding an amino acid or a cycloheximide-induced block in translation; (iii) Upf2 and Upf3X are detected on CBP80-bound mRNA but not eIF4E-bound mRNA; and (iv) mRNA immunopurified using anti-Upf3/3X antibody is bound by CBP80 but not eIF4E (Ishigaki *et al.*, 2001). These data indicate that NMD generally is confined to CBP80-bound mRNA, before CBP80 is replaced by eIF4E, and before Upf2 and Upf3X dissociate from the vicinity of splicing-generated exon-exon junctions (Ishigaki *et al.*, 2001).

We aimed to gain a broader understanding of the structural rearrangements that typify CBP80- and eIF4E-

bound transcripts as a consequence of pre-mRNA synthesis, pre-mRNA splicing, mRNA export and mRNA translation. Here, we report the characterization of proteins bound to spliced mRNA immunopurified from nuclear and cytoplasmic fractions of Cos cells using anti-CBP80, anti-eIF4E or anti-Upf3/3X antibody under conditions that preserve RNP. In nuclear fractions, anti-CBP80 antibody and anti-Upf3/3X antibody, but not anti-eIF4E antibody, immunopurified RNPS1, Y14, SRm160 and REF/Aly, all of which are components of the EJC. Also present in the anti-CBP80 and anti-Upf3/3X immunopurifications of nuclear fractions were the NMD factors Upf3X and Upf2 and the mRNA export factor TAP. RNase treatment demonstrated that immunopurification of RNPS1, Y14, SRm160, REF/Aly and Upf2, but not CBP80, using anti-Upf3/3X antibody was independent of RNA, consistent with these proteins constituting the EJC and associating with Upf3/3X. Furthermore, immunopurification of RNPS1, Y14, Upf3X, Upf2 and, to a lesser extent, SRm160 and REF/Aly using anti-CBP80 antibody was dependent on RNA, suggesting that SRm160 and REF/Aly, in addition to being part of the EJC, may interact with CBP80. The immunopurification of TAP using anti-Upf3/3X antibody was also dependent on RNA, suggesting that TAP has a relatively weak association with the EJC or is present in association with non-EJC mRNA-binding proteins, or both. All EJC components were detected in cytoplasmic fractions immunopurified using anti-CBP80 antibody in a way that is RNase sensitive, indicating that they are exported with mRNA from the nucleus to the cytoplasm before completely dissociating. Data demonstrating that Upf2 and Upf3X are exported to the cytoplasm in association with CBP80 and the EJC are consistent with the understanding that some mRNAs are subject to cytoplasmic NMD. The finding that eIF4E-bound transcripts that co-purify with nuclei are not bound by the EJC, together with the detection of significant levels of CBP80-bound pre-mRNA but not eIF4E-bound pre-mRNA, indicate that eIF4E replaces CBP80/CBP20 concomitant with or after dissociation of the EJC. In support of the idea that CBP80/CBP20 rather than eIF4E binds to nascent transcripts, antibody against the C-terminal domain of RNA polymerase II immunopurified CBP80 but not eIF4E.

Results

β -globin mRNA that co-purifies with nuclei is bound by CBP80 or eIF4E

CBP80 together with CBP20 comprise the major nuclear cap-binding complex that is added co-transcriptionally and functions in nuclear RNA processes such as pre-mRNA splicing and 3' end formation (Izaurralde *et al.*, 1994; Lewis and Izaurralde, 1997). Like CBP80 in *Chironomus tentans* (Visa *et al.*, 1996) and *S.cerevisiae* (Shen *et al.*, 2000), mammalian CBP80 is a nucleocytoplasmic shuttling protein (Ishigaki *et al.*, 2001). Given that the bulk of cellular translation involves mRNA bound by the major cytoplasmic cap-binding protein, eIF4E, it makes sense that eIF4E replaces CBP80 at some point after mRNA export to the cytoplasm (reviewed in Gingras *et al.*, 1999). However, the finding that a fraction of cellular eIF4E localizes to the nucleus (Lejbkovicz

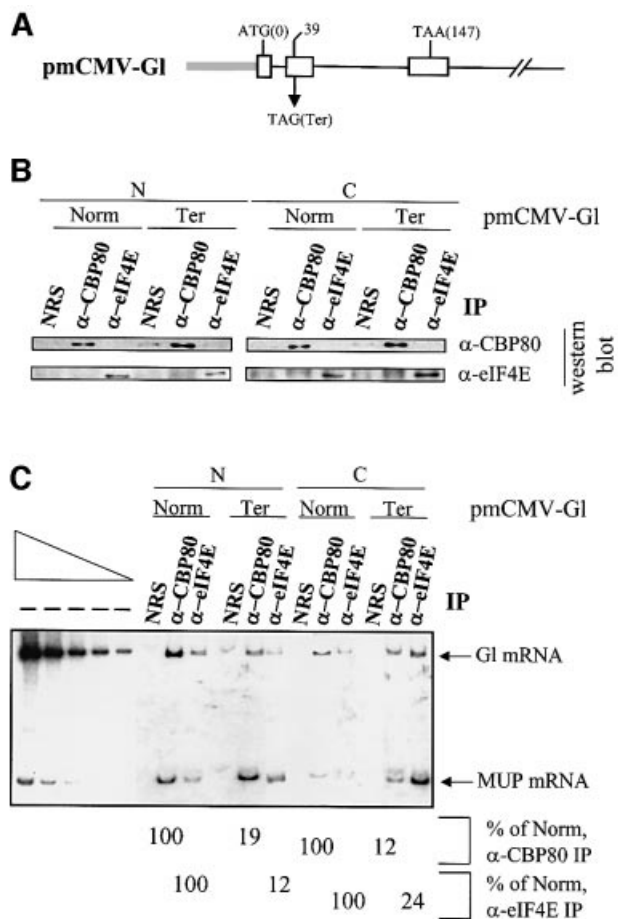


Fig. 1. CBP80 and eIF4E are bound to mRNA in both nuclear and cytoplasmic cell fractions. (A) Structure of the pmCMV-GI test plasmids, which harbor either a nonsense-free GI allele that terminates translation at codon 147 (Norm) or a GI allele carrying a TAG nonsense codon at position 39 (Ter). (B) Cos cells were transiently transfected with a test pmCMV-GI plasmid and the reference plasmid phCMV-MUP. Nuclear (N) and cytoplasmic (C) fractions were then immunopurified under conditions that preserved RNP using either normal rabbit serum (NRS), anti-CBP80 antibody (α -CBP80) or anti-eIF4E antibody (α -eIF4E), and subjected to western blotting using anti-CBP80 antibody or anti-eIF4E antibody. Nuclear fractions were free of cytoplasm as evidenced by the absence of detectable reactivity with anti-eIF4E antibody (data not shown). For easy visualization, bands corresponding to antibody are not shown. (C) The levels of GI and MUP mRNAs were quantitated in RNA prepared from each immunopurification using RT-PCR. The leftmost five lanes, which analyze decreasing amounts of RNA before immunopurification (-IP), demonstrate that the conditions of RT-PCR were quantitative. Numbers below the figure represent the level of GI mRNA normalized to the level of MUP mRNA, where normalized levels of nonsense-free (Norm) mRNA in each anti-CBP80 and anti-eIF4E immunopurification of each fraction are defined as 100%. Results are representative of two independently performed experiments where the efficiency of NMD varied by no more than 12%. The immunopurification using anti-eIF4E and cytoplasmic sample was less efficient for the transfection using pmCMV-GI Norm than for the transfection using pmCMV-GI Ter, explaining the reduced levels of eIF4E detected by western blotting (B) and RNA detected by RT-PCR (C).

et al., 1992; Dostie *et al.*, 2000) indicates that eIF4E could also have a nuclear function.

We first determined whether eIF4E that co-purifies with nuclei associates with mRNA. Cos cells were transiently transfected with two plasmids: a test pmCMV-GI plasmid

that produces β -globin (GI) mRNA (Zhang *et al.*, 1998; Figure 1A), and the reference phCMV-MUP plasmid that produces mRNA for the mouse major urinary protein (MUP; Belgrader and Maquat, 1994) and served to control for variations in the efficiencies of cell transfection and RNA recovery. After 40 h, cells were lysed, nuclear and cytoplasmic fractions were generated, and immunopurifications were performed under conditions that preserve RNP (Ishigaki *et al.*, 2001). Briefly, lysates first were cleared by incubation with protein A-agarose beads and subsequently incubated with rabbit anti-CBP80 antibody, rabbit anti-eIF4E antibody or, as a control for non-specific immunopurification, normal rabbit serum (NRS). After 90 min at 4°C, yeast RNA and protein A-agarose beads were added, and the incubation was continued for another 60 min. The beads were washed extensively, bound material was eluted with SDS-containing buffer, and protein and RNA were purified for analysis by western blotting and RT-PCR, respectively.

Results obtained using western blotting were consistent with previous findings (Ishigaki *et al.*, 2001): anti-CBP80 antibody immunopurified only CBP80 in both cellular fractions, anti-eIF4E antibody immunopurified only eIF4E in both cellular fractions; and the control NRS did not immunopurify either protein in either cell fraction (Figure 1B). RT-PCR demonstrated that each antibody immunopurified GI and MUP mRNAs and that, regardless of the cellular fraction or cap-binding protein, the level of nonsense-containing (Ter) GI mRNA was reduced to an average of $17 \pm 7\%$ the level of nonsense-free (Norm) mRNA (Figure 1C). Therefore, eIF4E is associated with mRNA that co-purifies with nuclei.

CBP80 but not eIF4E is detectably bound to intron-containing GI and SV40 T antigen pre-mRNAs

Since CBP80-bound but not eIF4E-bound mRNA was found to be associated with factors required for NMD, the reduction in the abundance of nonsense-containing eIF4E-bound mRNA was attributed to NMD that occurred prior to the exchange of CBP80 for eIF4E (Ishigaki *et al.*, 2001). Therefore, intron-containing pre-mRNA may be bound primarily by CBP80 rather than eIF4E. Alternatively, it may be bound by CBP80 or eIF4E, but the latter is a dead-end product that fails to be spliced and, therefore, fails to recruit Upf3/3X and Upf2 (Ishigaki *et al.*, 2001; see below). One way to determine whether eIF4E-bound mRNAs generally derive from CBP80-bound mRNAs or, alternatively, eIF4E binds to pre-mRNA is to identify the cap-binding protein on intron-containing RNA: if it is only CBP80, then eIF4E-bound RNA must derive from CBP80-bound RNA. To this end, the experiment described above was repeated. However, only nuclear fractions were prepared, and intron-containing GI pre-mRNA was analyzed instead of GI mRNA. Furthermore, the anti-eIF4E antibody derived from either rabbit or mouse. When mouse anti-eIF4E antibody was used, the level of non-specific immunopurification was controlled for using mouse IgG.

RT-PCR demonstrated that the levels of MUP mRNA immunopurified using either anti-CBP80 antibody or mouse anti-eIF4E antibody were higher than non-specific, i.e. higher than the levels immunopurified using NRS or mouse IgG, respectively (Figure 2A), consistent with

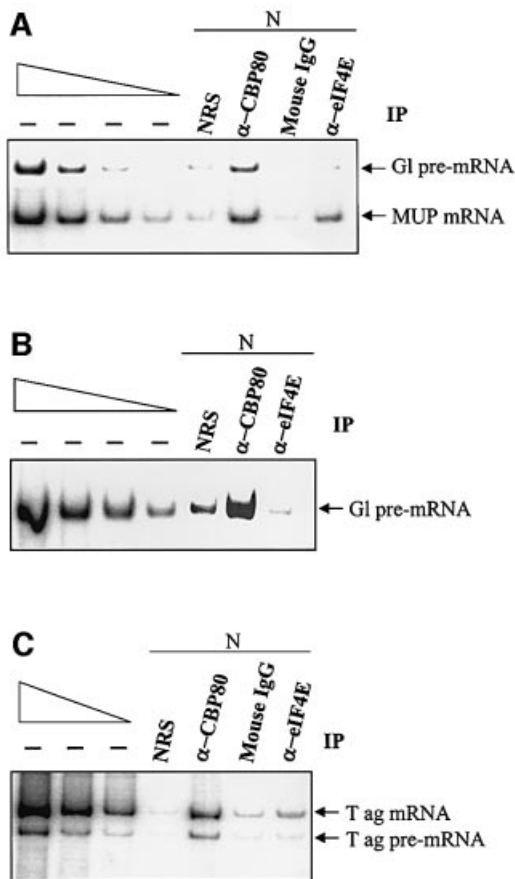


Fig. 2. CBP80 but not eIF4E is detectably bound to intron-containing pre-mRNA. (A) As in Figure 1, except that only nuclear fractions and Gl pre-mRNA rather than Gl mRNA were analyzed. Furthermore, mouse anti-eIF4E antibody was used instead of rabbit anti-eIF4E antibody and mouse IgG served as control for non-specific immunopurification using mouse anti-eIF4E antibody. (B) As in (A) except that rabbit anti-eIF4E antibody was used in place of mouse anti-eIF4E antibody and the analysis of MUP mRNA was omitted. (C) As in (A) except that SV40 pre-mRNA and mRNA were analyzed. Results are representative of other independently performed experiments, including those using rabbit anti-eIF4E antibody and NRS as a control. Taking the sum of Gl or SV40 pre-mRNA immunopurified by anti-CBP80 antibody and anti-eIF4E antibody in each panel as 100%, the amount of CBP80-bound pre-mRNA was $96 \pm 6\%$ and the amount of eIF4E-bound pre-mRNA was $4 \pm 6\%$.

previous findings (Ishigaki *et al.*, 2001; Figure 1). In contrast, while the level of Gl pre-mRNA immunopurified using anti-CBP80 antibody was higher than non-specific, the level of Gl pre-mRNA immunopurified using mouse anti-eIF4E antibody was barely above non-specific (Figure 2A). The essential failure to detect eIF4E-bound Gl pre-mRNA was reproduced using rabbit anti-eIF4E antibody (Figure 2B). Therefore, CBP80 is readily detected in association with Gl pre-mRNA whereas eIF4E is not.

To determine if this result can be extended to another pre-mRNA, the immunopurification was repeated using untransfected Cos cells. RT-PCR was then used to assay the relative levels of SV40 T antigen (ag) pre-mRNA and mRNA. The level of SV40 T ag mRNA immunopurified using either anti-CBP80 or mouse anti-eIF4E antibody was higher than non-specific (Figure 2C). However, as was

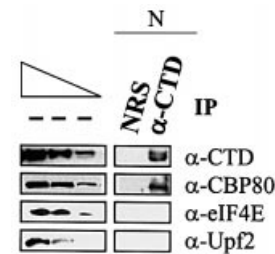


Fig. 3. CBP80 but neither eIF4E nor Upf2 immunopurifies with the CTD. The nuclear fraction of 32×10^6 untransfected Cos cells was immunopurified using either NRS or antibody to the C-terminal domain of RNA polymerase II (α -CTD), and protein from either 8×10^5 cells (CTD) or 32×10^5 cells (CBP80, eIF4E and Upf2) was subjected to western blotting using anti-CTD, anti-CBP80, anti-eIF4E or anti-Upf2 antibody. The three leftmost lanes, which analyze 2-fold dilutions of nuclear protein from 1.6×10^5 cells (CTD and CBP80) or 6.4×10^5 cells (eIF4E and Upf2) before immunopurification ($-IP$), demonstrate that the conditions of western blotting were semi-quantitative. The two forms of CTD probably differ in the degree of phosphorylation (Dubois *et al.*, 1994). Results are representative of 2–4 independently performed experiments, depending on the antibody used in western blotting.

the case for Gl pre-mRNA, the level of SV40 T ag pre-mRNA immunopurified using anti-CBP80 antibody was significantly higher than non-specific, while the level immunopurified using mouse anti-eIF4E antibody was only slightly higher than non-specific (Figure 2C).

We conclude that the vast majority of Gl and SV40 T ag pre-mRNAs is bound by CBP80 rather than eIF4E, suggesting that eIF4E-bound mRNA derives primarily from transcripts initially bound by CBP80.

CBP80 but not eIF4E co-immunopurifies with the C-terminal domain of RNA polymerase II

Given that 5' capping enzymes bind the phosphorylated C-terminal domain (CTD) of transcribing RNA polymerase II molecules (McCracken *et al.*, 1997; Schroeder *et al.*, 2000; Pei *et al.*, 2001), and since proteins known to function in pre-mRNA splicing and 3'-end formation also associate with the CTD (reviewed in Hirose and Manley, 2000; Proudfoot, 2000), an additional way to determine whether CBP80-bound mRNA is generally a precursor to eIF4E-bound mRNA might be to determine whether CBP80 or eIF4E also co-immunopurify with the CTD. To test this idea, the nuclear fraction of untransfected Cos cells was immunopurified using anti-CTD antibody, and the presence of CBP80 and eIF4E was tested.

The results of western blotting indicate that anti-CTD antibody immunopurified CTD, as expected, and, remarkably, CBP80 but not eIF4E (Figure 3; data not shown for the analysis of larger amounts of immunopurified material). This finding is consistent with the finding that intron-containing pre-mRNA is bound by CBP80 rather than eIF4E (Figure 2) and offers additional support for a precursor-product relationship between CBP80-bound mRNA and eIF4E-bound mRNA. Another protein that failed to immunopurify with the CTD was Upf2 (Figure 3), which was chosen as a negative control based on its cytoplasmic localization (Lykke-Andersen *et al.*, 2000; Serin *et al.*, 2001).

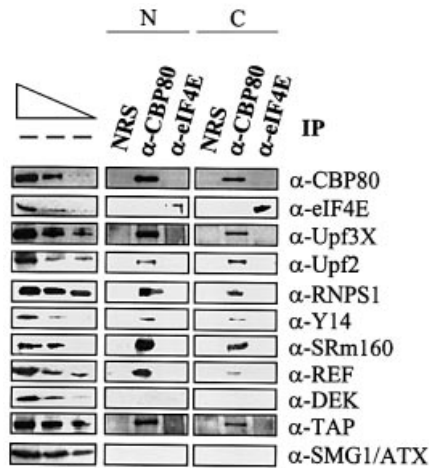


Fig. 4. Nuclear and cytoplasmic CBP80, unlike nuclear and cytoplasmic eIF4E, co-immunopurify with RNPS1, Y14, SRm160, REF/Aly, TAP, Upf3X and Upf2. Nuclear and cytoplasmic fractions of 32×10^6 untransfected Cos cells were immunopurified using anti-CBP80 antibody, anti-eIF4E antibody or, as a control, NRS. Immunopurified protein from 8×10^5 or 32×10^5 cells was then analyzed by western blotting using the antibodies specified. The three leftmost lanes, which analyze 2-fold dilutions of nuclear protein from 8×10^5 or 32×10^5 cells before immunopurification (– IP), demonstrate that the conditions of western blotting were semi-quantitative. Notably, Upf3 probably co-migrates with the heavy chain of anti-hUpf3/3X antibody and, therefore, was not assayable.

Anti-CBP80 antibody, unlike anti-eIF4E antibody, co-immunopurifies with RNPS1, Y14, SRm160, REF/Aly, TAP, Upf2 and Upf3X in nuclear and cytoplasmic fractions

To gain insight into the dynamics of mRNP remodeling before and after eIF4E replaces CBP80 at the mRNA cap, the immunopurifications were repeated using nuclear and cytoplasmic fractions from untransfected Cos cells and analyzed for a variety of proteins using western blotting. Initially, RNPS1, Y14, SRm160, REF/Aly and DEK, which are components of the mRNA EJC that is deposited as a consequence of pre-mRNA splicing *in vitro*, were analyzed. Remarkably, all components except DEK were detected in immunopurifications of both nuclear and cytoplasmic fractions using anti-CBP80 antibody (Figure 4). The failure to detect DEK suggests that either it is not present in the complex *in vivo* or that conditions used in the immunopurification removed DEK. A similar conclusion was drawn from immunopurifications using unfractionated mammalian cells expressing FLAG-tagged proteins, where all tagged components of the EJC were detected except for DEK (Lykke-Andersen *et al.*, 2001). Our results indicate that RNPS1, Y14, SRm160 and REF shuttle to the cytoplasm, presumably in association with spliced mRNA (see below). Considering that SRm160 does not shuttle between the two nuclei of mammalian cell heterokaryons (Lykke-Andersen *et al.*, 2001; Y.Ishigaki, B.Blencowe and L.E.Maquat, unpublished data), detection of SRm160 in the cytoplasmic fraction by immunopurification using anti-CBP80 antibody suggests that SRm160 may have a limited trajectory in the cytoplasm. Consistent with our failure to detect eIF4E-bound pre-mRNA, no components of the EJC were detected in either cell fraction using anti-eIF4E antibody (Figure 4).

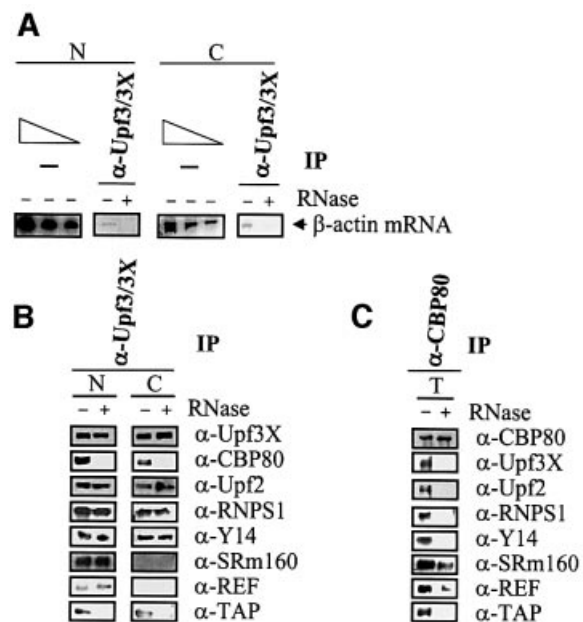


Fig. 5. RNPS1, Y14, SRm160, REF/Aly and Upf2 co-immunopurify with Upf3X in an RNase-insensitive manner, while RNPS1, Y14, REF/Aly, TAP, Upf3X and Upf2 co-immunopurify with CBP80 in an RNase-sensitive manner. As in Figure 4 except that an additional immunopurification using anti-Upf3/3X was performed, total (T) lysate was immunopurified using anti-CBP80 antibody, RNase was added to half of each sample prior to immunopurification, RNA as well as protein were prepared, and western blot analyses of eIF4E and DEK were omitted. (A) RT-PCR quantitation of the level of β -actin mRNA before or after immunopurification using anti-Upf3/3X antibody in order to demonstrate the efficiency of RNase treatment. Prior to immunopurification, half of the sample was treated with RNase (+), and the other half was not treated with RNase (–). (B) Western blot analysis of protein immunopurified using anti-Upf3/3X antibody from nuclear and cytoplasmic fractions. (C) Western blot analysis of protein immunopurified using anti-CBP80 antibody from total cell extract.

The mRNA export factor TAP and the NMD factors Upf2 and Upf3X were also detected in anti-CBP80 antibody immunopurifications of nuclear and cytoplasmic fractions but not anti-eIF4E immunopurifications of either fraction (Figure 4). The failure to detect either Upf protein using anti-eIF4E antibody is consistent with data demonstrating that eIF4E-bound mRNA is largely immune to NMD (Ishigaki *et al.*, 2001). SMG1/ATX, the PIK-related protein kinase thought to be required for NMD because of its role in phosphorylating Upf1 (Denning *et al.*, 2001; Yamashita *et al.*, 2001; K.M.Brumbaugh, D.M.Otterness, X.Li, F.Lejeune, R.S.Tibbetts, L.E.Maquat and R.T.Abraham, unpublished data), was not detected in either anti-CBP80 or anti-eIF4E immunopurification (Figure 4). This suggests that, like Upf1, SMG1/ATX is not a stable component of mRNP, at least under the purification conditions used here.

Evidence that RNPS1, Y14, SRm160, REF/Aly, TAP, Upf2 and Upf3X are components of or associate with the EJC on CBP80-bound mRNA in mammalian cells

In order to determine if RNPS1, Y14, SRm160 and REF/Aly immunopurify with anti-CBP80 antibody as components of the EJC of mRNP, the immunopurifications were repeated using anti-Upf3/3X antibody or

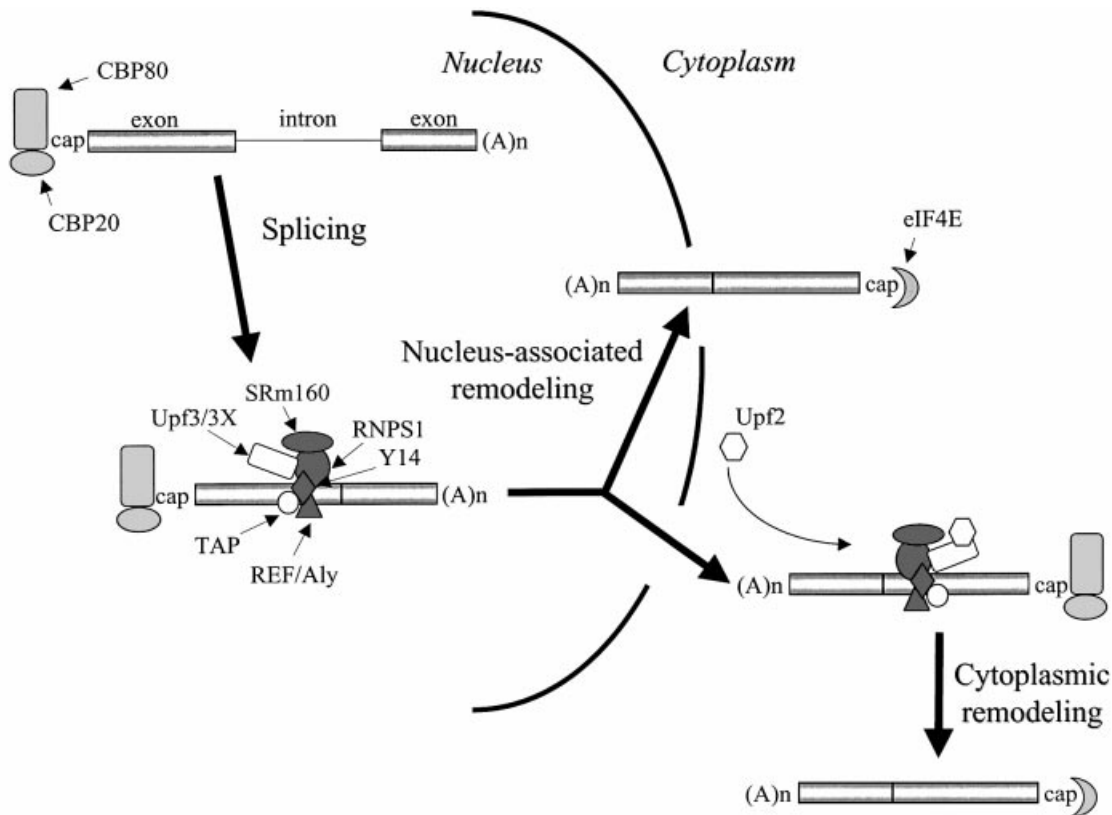


Fig. 6. Model for the dynamics of mRNP structure in mammalian cells as a consequence of pre-mRNA capping, pre-mRNA splicing and mRNA export. In the nucleus, CBP80-bound pre-mRNA is spliced to generate CBP80-bound mRNA. This mRNA is bound by an EJC (dark gray shapes), consisting of RNPS1, Y14, SRm160 and REF/Aly, located 20–24 nucleotides upstream of the exon–exon junction. The EJC recruits the NMD factor Upf3/3X and the mRNA export factor TAP. Some mRNPs (possibly those subject to nucleus-associated NMD) undergo remodeling while associated with the nucleus (nucleus-associated remodeling). Other mRNPs (including those subject to cytoplasmic NMD) undergo remodeling after export to the cytoplasm (cytoplasmic remodeling). In both types of remodeling, CBP80 is replaced by eIF4E and the EJC is lost. Data indicate that the mostly perinuclear Upf2 protein is recruited by the EJC at a point when mRNA co-purifies with nuclei. In theory, recruitment could occur during mRNA export. However, if nucleus-associated remodeling takes place in the nucleoplasm, it is conceivable that an as yet undetected pool of Upf2 is nucleoplasmic. The single exon–exon junction exemplified is likely to typify all splicing-generated exon–exon junctions, regardless of position relative to the mRNA 3' end.

anti-CBP80 antibody either with or without the addition of RNase prior to the immunopurification step. Data demonstrating that Upf3/3X is recruited to mRNP by an EJC formed *in vitro* in mammalian cell extract (Kim *et al.*, 2001a) or *in vivo* in intact mammalian cells using FLAG-tagged Upf3 produced by transient transfection (Lykke-Andersen *et al.*, 2001) indicate that the EJC and CBP80 bind to distinct regions of mRNA. Therefore, components of or proteins recruited by the EJC would be expected to immunopurify with anti-Upf3/3X antibody both before and after RNase treatment, but with anti-CBP80 antibody only before and not after RNase treatment.

RNase treatment was shown to be effective by the disappearance of β -actin mRNA in immunopurified samples after exposure to RNase (Figure 5A; data not shown for the immunopurification using anti-CBP80 antibody). As expected, anti-Upf3/3X antibody immunopurified Upf3X in nuclear fractions with or without exposure to RNase (Figure 5B) and anti-CBP80 antibody immunopurified CBP80 in total cell extract with or without exposure to RNase (Figure 5C). Furthermore, anti-Upf3/3X antibody immunopurified CBP80 in both fractions only in the

absence of RNase, but immunopurified Upf2, RNPS1 and Y14 in both fractions in the absence or presence of RNase (Figure 5B). These results, together with the finding that anti-CBP80 antibody immunopurified Upf3X, Upf2, RNPS1 and Y14 in total cell extract only in the absence of RNase (Figure 5C), indicate that Upf3X and Upf2 form a complex with the EJC that is conserved after mRNA export to the cytoplasm. SRm160 and REF/Aly immunopurified with anti-Upf3/3X antibody in the nuclear fraction with or without exposure to RNase (Figure 5B), as would be expected of EJC components. However, they failed to immunopurify with anti-Upf3/3X antibody in the cytoplasmic fraction (Figure 5B), and they immunopurified with anti-CBP80 antibody in a partially RNase-insensitive manner (Figure 5C). Considering that both proteins immunopurify with anti-CBP80 antibody in the cytoplasmic fraction (Figure 4), they may undergo a remodeling in the cytoplasm that provides a more direct connection to CBP80. The finding that TAP immunopurifies with anti-Upf3/3X antibody and anti-CBP80 antibody in an RNase-sensitive manner in nuclear and cytoplasmic fractions (Figure 5B and C) does not preclude the possibility that it is recruited by the EJC. In fact, TAP could be only loosely

associated with the EJC, making it susceptible to removal by RNase treatment. However, it is also possible that some TAP is recruited by the EJC and the majority is recruited by non-EJC mRNA-binding proteins (see Discussion).

Discussion

This study aimed to characterize proteins associated with CBP80-bound and eIF4E-bound RNA in mammalian cells before and after export from the nucleus to the cytoplasm. Notably, endogenous proteins rather than proteins produced from introduced genes were assayed in order to avoid generating artifactual interactions attributable to expression at abnormal levels. Even though both CBP80 and eIF4E are present in nuclei (Ohno *et al.*, 1990; Izaurralde *et al.*, 1992; Lejbkiewicz *et al.*, 1992; Dostie *et al.*, 2000) and eIF4E is detected in association with mRNA that co-purifies with nuclei (Figure 1), only CBP80 is readily detected in association with intron-containing pre-mRNA (Figure 2). The failure to detect appreciable amounts of eIF4E-bound pre-mRNA is consistent with data demonstrating that immunodepletion of CBP80 from HeLa cell extracts strongly reduces the efficiency of pre-mRNA splicing (Izaurralde *et al.*, 1994). It is also consistent with the detection of CBP80 but not eIF4E in association with the CTD (Figure 3). We find that CBP80 and not eIF4E co-immunopurifies with components of the EJC (Figure 4). In verification that our conditions allow for the isolation of mRNP and, therefore, a characterization of its structure, we also demonstrate that immunopurification of components of the EJC using anti-CBP80 antibody is sensitive to RNase whereas immunopurification of these components using anti-Upf3/3X antibody is not (Figure 5). This result is expected of cap-bound CBP80 and distally bound components of the EJC, the latter of which recruits Upf3/3X. Our findings are consistent with data demonstrating that either FLAG-Upf3 or FLAG-Upf3X expressed transiently in 293T cells immunopurifies Y14 and REF/Aly independently of RNA (Kim *et al.*, 2001b; RNPS1, SRm160 and Upf2 were not tested and, in contrast to our results, only Flag-Upf3 immunopurified TAP in a manner that was partially dependent on RNA). We conclude that eIF4E binds RNA after splicing and concomitant with or after removal of the EJC.

Given that CBP80-bound mRNA is the substrate for NMD and the NMD of some mRNA takes place in the cytoplasm, the replacement of CBP80 by eIF4E on mRNA subject to cytoplasmic NMD must take place in the cytoplasm at least some of the time (Ishigaki *et al.*, 2001; Figure 6). However, the presence of eIF4E-bound mRNA in the nuclear fraction (Figure 1) indicates that the replacement of CBP80 by eIF4E on mRNA also takes place before mRNA release into the cytoplasm at least some of the time (Figure 6). In fact, replacement before release might typify mRNAs subject to nucleus-associated NMD. In theory, replacement before release could take place during transit across the nuclear pore, at a point when mRNA would co-purify with nuclei but have access to cytoplasmic ribosomes. In support of this possibility, mRNP in the insect *C.tentans* has been shown using electron microscopy to exit the nucleus 5' end-first and associate with cytoplasmic ribosomes before its 3' end transits the nuclear pore complex (Mehlin *et al.*, 1992;

Daneholt, 1997). Alternatively, since the EJC is not required for export (Huang and Steitz, 2001; Rodrigues *et al.*, 2001), replacement could take place in the nucleoplasm.

Our data are consistent with a growing number of studies demonstrating that mRNP undergoes significant remodeling, in keeping with the requirements of mRNA export and NMD (Lou and Reed, 1999; Lou *et al.*, 2001; Kataoka *et al.*, 2000; Le Hir *et al.*, 2000b, 2001a; Zhou *et al.*, 2000; Kim *et al.*, 2001a,b). RNPS1, Y14, SRm160 and REF/Aly, which are components of the EJC, and Upf2, Upf3X and TAP, which are recruited by the EJC either directly or indirectly (Le Hir *et al.*, 2000a, 2001a,b; Ishigaki *et al.*, 2001; Kim *et al.*, 2001b; Lykke-Andersen *et al.*, 2001), are detected on CBP80-bound mRNA in both nuclear and cytoplasmic fractions (Figure 4). Therefore, each protein first binds directly or indirectly to mRNA either in the nucleoplasm or during mRNA export, and at least a fraction of each protein remains bound after export to the cytoplasm.

Studies of *Xenopus* oocytes indicate that Y14 and SRm160 remain associated with spliced mRNA after export, while RNPS1, REF/Aly and TAP fail to be detected on exported mRNA (Le Hir *et al.*, 2001a). Differences in the behavior of RNPS1, REF/Aly and TAP between mammalian cells and *Xenopus* oocytes are not yet understood. The finding that these proteins are complexed with Y14, which remains bound to mRNA after export in *Xenopus* oocytes, suggests that there are bona fide differences between the fate of EJC components in mammalian cells and *Xenopus* oocytes. As another difference, Upf2 and Upf3X are associated with the EJC in both nuclear and cytoplasmic fractions of mammalian cells (Figures 4 and 5), whereas Upf3X is detected on mRNA only in *Xenopus* oocyte nuclei and Upf2 is detected on mRNA only in *Xenopus* oocyte cytoplasm (Le Hir *et al.*, 2001a). Unlike mammalian cells, *Xenopus* oocytes do not support NMD (J.Zhang, E.Lund and L.E.Maquat, unpublished data), which may reflect the observed differences in mRNP.

Very recently, UAP56 and epitope-tagged Mago have also been shown to associate with the EJC. UAP56, a DEAD-box helicase and splicing factor, and Sub2p, its ortholog in *S.cerevisiae*, are linked to nuclear mRNA export via interactions with REF/Aly and Yra1p, respectively (Lou *et al.*, 2001; Sträßer and Hurt, 2001). Mago joins the EJC by interacting with Y14 and has been shown to escort spliced mRNAs to the cytoplasm in *Xenopus* oocytes (Kataoka *et al.*, 2001; Le Hir *et al.*, 2001b). These studies join the growing number of observations that splicing, transport and translation are mechanistically linked through numerous steps of mRNP remodeling.

Data indicate that EJCs function in mRNA export (Lou and Reed, 1999; Zhou *et al.*, 2000; Kim *et al.*, 2001b; Le Hir *et al.*, 2001a; reviewed in Reed and Magni, 2001; Reed and Hurt, 2002). However, EJCs are not the only means by which mRNA is exported, as exemplified by the export of intronless RNAs in pathways that involve the nucleocytoplasmic shuttling proteins SRp20 and 9G8 (Huang and Steitz, 2001) and REF/Aly (Rodrigues *et al.*, 2001). Given that spliced and unspliced mRNAs use TAP to reach the cytoplasm (Rodrigues *et al.*, 2001), and considering that SRp20 and 9G8 have been shown to

cross-link to polyadenylated RNA in both nuclear and cytoplasmic cell fractions (Huang and Steitz, 2001), it is conceivable that spliced mRNAs are bound by non-EJC proteins that recruit TAP for the purpose of export, possibly explaining at least in part the RNase sensitivity of TAP co-immunopurification with Upf3/3X (Figure 5B). Along similar lines, T-cell receptor- β mRNA was shown recently to harbor a novel *cis*-acting sequence distinct from an EJC that triggers NMD, presumably by recruiting Upf factors (Wang *et al.*, 2002). Future studies that continue to characterize the nature of spliced mRNP should lend important insight into effectors of mRNA synthesis, transport and half-life.

Materials and methods

Cell culture and transfections

Monkey kidney Cos-7 cells were cultured and, where indicated, transfected as previously described (Ishigaki *et al.*, 2001).

Cell fractionation, lysis and immunopurifications

Whole cells or nuclear and cytoplasmic fractions of cells ($3\text{--}4 \times 10^7$) were processed as previously described (Ishigaki *et al.*, 2001). For all immunopurifications except those using anti-CTD antibody, samples were first cleared by incubation with end-over-end rotation in the presence of 50 μ l of protein A-agarose beads (Boehringer Mannheim) in NET-2 buffer, 1 mM phenylmethylsulfonyl fluoride (PMSF; Sigma), 2 mM benzamidine (Sigma) and, when RNase was not used, 100 U of RNase inhibitor (Promega) for 30 min at 4°C followed by centrifugation at 10 000 g. Cleared samples were then rotated in the presence of either normal rabbit antiserum (NRS; Gibco-BRL) or mouse IgG (Sigma), anti-CBP80 antibody (Izaurralde *et al.*, 1994), anti-eIF4E antibody (Morley and McKendrick, 1997; Santa Cruz Biotechnology) or anti-hUpf3/3X antibody (Ishigaki *et al.*, 2001) for 90 min at 4°C, after which 2 mg of yeast RNA (Sigma) and protein A-agarose beads were added. The incubation was continued for another 60 min at 4°C. The beads were washed 5–6 times with NET-2 buffer, suspended in 50 μ l of SDS sample buffer (Ishigaki *et al.*, 2001) that either did or, for the analysis of RNPS1 and Upf3 that co-migrated with antibody, did not contain 12% 2-mercaptoethanol, and split into two portions. In some experiments, RNase A (10 μ g; Sigma) or, as a control, bovine serum albumin (BSA; 10 μ g; New England Biolabs) was added to the beads and incubated for 30 min at 37°C before elution using SDS sample buffer. One portion (three-fifths) was used as a source of protein, which was analyzed by SDS-PAGE and western blotting. The other portion (two-fifths) was extracted with phenol, chloroform and isoamyl alcohol, and precipitated using ethanol. Precipitates were dissolved in 20 μ l of RQ1 DNase buffer (Promega), treated with 10 U of DNase I (Promega) for 30 min at 37°C, extracted, precipitated, dissolved in 20 μ l of water, and used as a source of RNA for RT-PCR (Sun *et al.*, 1998).

For immunopurifications using anti-CTD antibody and, as a control, NRS, the concentration of NaCl in NET-2 buffer was decreased from 300 to 50 mM.

Western blotting

Protein in immunopurifications (1–5 μ l) was electrophoresed in 7.5% (for SMG1/ATX), 10% (for CBP80, Upf2, SRm160, TAP and CTD) or 12% (for eIF4E, Y14, REF, DEK, Upf3X and RNPS1) polyacrylamide. Protein was transferred to Hybond ECL nitrocellulose (Amersham), and probed with antibody against CBP80 (a gift from E.Izaurralde), eIF4E (a gift from S.Morley), Upf3/3X (Serin *et al.*, 2001), Upf2 (Serin *et al.*, 2001), RNPS1 (a gift from A.Mayed), Y14 (a gift from G.Dreyfuss), SRm160 (a gift from B.J.Blencowe), DEK (a gift from G.Grosveld), REF/Aly (a gift from E.Izaurralde), TAP (a gift from E.Izaurralde), SMG1/ATX (a gift from R.T.Abraham) or CTD (a gift from D.Bentley). In the case of cellular fractionations, nuclear fractions were deemed free of cytoplasm by the absence of detectable reactivity with antibody against eIF4A (a gift from S.Morely; Lejbkovicz *et al.*, 1992) and anti-PLC γ (a gift from R.T.Abraham). Reactivity to each primary antibody was detected using a 1:5000 dilution of anti-rabbit Ig, horseradish peroxidase (HRP)-linked whole antibody (Amersham; to detect CBP80, eIF4E, RNPS1, Y14, REF/Aly, TAP, Upf2, Upf3X, SMG1/ATX and CTD), anti-mouse Ig, HRP-linked whole antibody (Amersham; to detect eIF4E), or anti-mouse IgM

peroxidase conjugate (Sigma; to detect SRm160). Reactivity of the secondary antibody was visualized by SuperSignal West Pico or West Femto Solution (Pierce).

RT-PCR

GI or MUP mRNAs in immunopurifications (10 μ l) were analyzed by RT-PCR as described (Ishigaki *et al.*, 2001). Alternatively, GI pre-mRNA was amplified using primers 5' GCCTATTGGTCTATTTCCC 3' (sense) and 5' CCTGAAGTTCTCAGGATC 3' (antisense). To assay for RNase activity, endogenous Cos-cell β -actin mRNA was amplified using 5' ATCTGGCACACACCTTCTACAATGAGCTGCG 3' (sense) and 5' CGTCATACTCTGCTTGCTGATCCACATCTGC 3' (antisense). Endogenous Cos cell SV40 T ag pre-mRNA and mRNA were amplified using primers 5' TGCAAGGAGTTTCATCCTG 3' (sense) and, respectively, 5' AGAATCAGTAGTTTAACACAC 3' and 5' TGAGCA-TAGTTATTAATAGCAG 3' (antisense), in which case the annealing reaction was performed at 50°C for 40 s and a total of 23 cycles. The simultaneous analysis of serial dilutions of RNA ensured that RT-PCR was quantitative. RT-PCR products were quantitated by PhosphorImaging (Molecular Dynamics).

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