RNA-binding proteins as developmental regulators

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Many RNA-binding proteins of the nucleus and cytoplasm, including pre-mRNA-, mRNA-, snRNA-, and pre-rRNA-binding proteins, contain a putative RNA-binding domain of approximately 90 amino acids whose amino acid sequence is conserved from yeast to man. The most highly conserved motif within this RNA-binding domain is an octapeptide, termed the ribonucleoprotein consensus sequence [RNP-CS], which is an identifying feature of this group of proteins. Frequently, these proteins contain several similar, but nonidentical, RNP-CS-type RNA-binding domains. All of these proteins also contain at least one auxiliary domain that is unique to each type of protein and most likely functions in protein-protein interactions. Many, if not all, of the RNP-CS-type proteins display binding preferences for specific RNA sequences, and several have been shown to interact with pre-mRNA sequences important for pre-mRNA processing. Recent work has shown that the proteins encoded by several developmental loci in Drosophila and maize contain RNP-CS and, therefore, are most likely RNA-binding proteins. Here we provide an overview of the structural characteristics of these proteins and speculate on how the modular structure of RNP-CS-type RNA-binding proteins may facilitate their participation in pathways that regulate development at the post-transcriptional level.

Identification of the RNP consensus sequence

The RNP-CS [Adam et al. 1986; Swanson et al. 1987] was identified initially in the yeast mRNA poly[A]-binding protein [PABP] [Adam et al. 1986; Sachs et al. 1986] and in the mammalian heterogeneous nuclear ribonucleoprotein particle [hnRNP] protein A1 [Cobianchi et al. 1986, Riva et al. 1986], and subsequently has been found in many other proteins [Lahiri and Thomas 1986; Theissen et al. 1986, Grange et al. 1987, Haynes et al. 1987, Jong et al. 1987; Lapeyre et al. 1987; Sillikens et al. 1987, Amrein et al. 1988, Bell et al. 1988; Chambers et al. 1988, Deutscher et al. 1988, Gomez et al. 1988, Preugschat and Wold 1988, Robinow et al. 1988; Goralski et al. 1989]. This octapeptide consensus sequence, Lys/Arg-Gly-Phe-Tyr-Gly/Ala-Phe-Val-X-Phe/Tyr, is the most highly conserved segment in a generally conserved region consisting of about 90 amino acids present at least once in all of these proteins. This region, because of its general conservation as a unit and because of experimental findings discussed below, is considered to be an RNA-binding domain. In this review, we shall refer to the 90-amino-acid region as an RNP-CS-type RNA-binding domain and to the more highly conserved octapeptide sequence as RNP-CS.

Figure 1 lists the proteins in which the RNP-CS sequence has been found thus far. All of the RNP-CS-type proteins are known RNA-binding proteins, except for the developmental control genes discussed below. The RNP-CS motif defines only one class of RNA-binding protein, because many well-characterized RNA-binding proteins exist [e.g., ribosomal proteins, viral nucleocapsid proteins] that do not contain this motif. In addition to the RNP-CS [RNP 1 in Fig. 1], alignment of the relevant RNP-CS-containing segments of each of the proteins reveals that several other amino acids at specific positions in the RNP-CS-type RNA-binding domain also are conserved, for example, a 6-amino-acid segment, RNP 2 [Dreyfuss et al. 1988], located about 30 amino acids amino-terminal to the RNP-CS. RNP 2 is less well conserved than the RNP-CS, yet the general character of its amino acids is preserved. Sequence similarities also have been reported between the eukaryotic RNP-CS and several prokaryotic ssDNA-binding proteins [including T4 gp32, M13 gp5, and E. coli ssb] and bacterial σ transcription factors [Helmann and Chamberlin 1988].

What is the evidence that the RNP-CS-type RNA-binding domain is responsible for the RNA-binding properties displayed by each of these proteins? Characterization of UP1, the mammalian single-stranded DNA-binding protein [Herrick and Alberts 1976], has revealed that this 24-kD polypeptide is an amino-terminal fragment of the hnRNP protein A1 that results from proteolysis in vitro [Kumar et al. 1986, Riva et al. 1986]. UP1, which is composed of two similar domains of approximately 90 amino acids each, retains the ability to bind single-stranded DNA or RNA [Kumar et al. 1986; Riva et al. 1986], and recent photo-cross-linking experiments have demonstrated that residues of the RNP-CS and RNP 2 within each of these two domains is directly in contact with RNA [Merrill et al. 1988]. Furthermore, experiments using fragments of the human hnRNP C1 protein that contain only the first 150 amino-terminal
Modular structure of RNP-CS-type proteins

Several RNP-CS-type RNA-binding domains often occur in an individual protein (Fig. 1) and usually are located in the amino-terminal portion of the protein. Examination of these repeats in corresponding proteins from divergent organisms, especially with regard to evolutionary relationships, has implications for the possible activities of these proteins. Comparison of the RNA-binding domains within a single protein containing multiple RNA-binding domains typically shows only 30% amino acid identity. However, comparison of the corresponding domain in the same protein from divergent species (e.g., alignment of domain 1 from yeast and mammalian PABPs) shows 70% identity. The contribution of particular residues in the RNA-binding domain is required for binding (Sachs et al. 1987). Nevertheless, the RNP-CS is a strong indicator of the specific RNA-binding properties of the native protein (C. Burd, E. Mortenson, and G. Dreyfuss, unpublished). Taken together, these data indicate that the RNP-CS and RNP 2 segments are integral components of an RNA-binding domain. The contribution of particular residues in the RNA-binding domain to RNA binding is not yet known. It is even possible that in some cases only part of the RNA-binding domain is required for binding (Sachs et al. 1987). Nevertheless, the RNP-CS is a strong indicator of RNA-binding function.

## Table: RNA Binding Domain consensus

| Protein        | Domain | mRNA | RNA Binding Domain |
|----------------|--------|------|--------------------|
| hnRNPs         |        |      |                    |
| A1 rat         | 1      | SPKEPEIQIRK | LGFQLG . SFTEDDES.LASHEFQ , NGTVDCVDMOPDRKPS . |
| Drosophila     | 1      | SITEPEEIMIRK | LGFRLG . DRYEDGE.NAHEKFXE , NGTVDCVDMOPDRKPS . |
| A1 rat         | 2      | RGPAHLTVKMK | LGFQLG . KEDTEKHEA.DGPLYQ . YINIVEXT . MDRGUS . |
| Drosophila     | 2      | SPKAGAVTVKMK | LGFQLG . KEDTHESLYQ , YIKTVYIVEXTIV . MDFGUS . |
| C1 human       |        | KTOPDRMRSMR | LFVICNL . NTIVVKKSDVERAF , YIKTVYIVEXTIV . CCSVH . |
| C1 xenopus     |        | KTOPDRMRSMR | YDFVCNL . NTIVVKKSDVERAF , YIKTVYIVEXTIV . CCSVH . |
| E(U2)human     |        | AMKTRPEVKMM | LGFQLG . SFTDTEP , IREYFQG , YGEVEKMDPNLEK . |

**Figure 1.** The 90-amino-acid RNA binding domains of RNP-CS-type proteins. The most highly conserved segments of these domains are denoted RNP 1 [RNP-CS] and RNP 2 and are in boldface type, as are other conserved amino acid residues located throughout the domain.

Conserved positions listed under RNP consensus, which were detected by pair-wise comparisons using the UWCGG GAP program and further refined by visual inspection, were selected by having no more than two different amino acids in each position in at least 20 out of the 32 domains; only one amino acid appears under RNP consensus when it alone fulfills this criterion. Protein sequences were obtained from the following references: hnRNP A1 from rat (Cobianchi et al. 1986) and protein (C. Burd, E. Mortenson, and G. Dreyfuss, unpublished). Nucleolin, indicate that this domain is sufficient to re...

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man] demonstrates strong amino acid similarities, typically greater than 60% identity [Dreyfuss et al. 1988]. Even more striking, the corresponding domains of the yeast and human PABPs contain segments of 8–10 amino acids that are completely identical [see pairwise comparison, Fig. 4 in Dreyfuss et al. 1988]. These segments are characteristic of the individual RNA-binding domain alone, and they are most likely indicative of functional specialization. In the A1 protein from mammals [rat] and Drosophila, domain 1 has an amino acid segment downstream of the RNP-CS in which 15 out of 16 amino acids are identical [Fig. 1, underlined region]; overall these corresponding domains share a 65% identity [Dreyfuss et al. 1988]. A schematic model depicting the modular assembly of RNP-CS-type proteins is presented in Figure 2. The patterns of evolutionary conservation among these domains suggest that they evolved by duplication and diversification of a common, ancestral RNA-binding protein gene bearing the prototypical RNP-CS. Subsequently, each RNA-binding domain, for example in the PABP or in the hnRNP protein A1, evolved independently. The diversification of the RNA-binding domains is particularly evident in isoforms of the A1 protein, where point mutations cause conservative amino acid substitutions in the first RNA-binding domain of this protein [Buvoli et al. 1988].

The remainder of a RNP-CS-type protein is characterized typically by a distinctive auxiliary domain(s). For example, the auxiliary domain is glycine-rich in the hnRNP protein A1, proline-rich in the polyadenylate binding protein [PABP], and glutamic and aspartic acid-rich in the hnRNP C proteins. The abundant nucleolar pre-rRNA-binding protein, nucleolin, has a glycine-rich domain similar to that of the hnRNP proteins A1 and A2 [Kumar et al. 1986] and an acidic domain similar to that of the hnRNP C proteins [Lapeyre et al. 1987]. These auxiliary domains appear likely to function primarily in protein–protein interactions, since all of these proteins are components of multiprotein ribonucleoprotein complexes.

**Figure 2.** Schematic model of the evolution and modular assembly of RNP-CS-type RNA-binding proteins. It is postulated that a prototype RNP-CS protein gene, encoding the ancestral RNA-binding domain protein, was subject to duplication and divergence, thereby expanding the range of RNA-binding specificities of RNP-CS-type proteins. The individual character of the diverged RNA-binding domains is represented by slanted lines or by stippled areas, whereas the conserved RNP-CS (RNP 1) is shown by the black box and RNP 2 by a smaller slashed box. In addition, auxiliary domains, composed of a distinctive set of amino acids, contribute to the domain character of RNP-CS-type proteins.
RNA-binding specificity
Several of the proteins listed in Figure 1 have binding specificity for certain RNA sequences. For example, the PABP binds preferentially to the poly[A] segments of mRNA. Recently, several of the hnRNP proteins, A1, C, and D have been shown to have RNA-binding specificity. The amino acid sequence of the hnRNP D proteins has not been determined yet, but the hnRNP A1 and C proteins, as well as the mRNP PABP, contain RNP-CS-type RNA-binding domains. Experimental efforts to purify individual hnRNP proteins led to the observation that several of these proteins display very high binding affinities for specific ribonucleotide homopolymers (Swanson and Dreyfuss 1988a). For example, the hnRNP C proteins have a striking affinity for poly[U], since they remain bound to this homopolymer at high salt concentrations (2 M NaCl). As reference, the yeast cytoplasmic PABP remains selectively bound to poly[A] in vitro only up to 1 M NaCl.

The finding that hnRNP C proteins are poly[U]-binding proteins led to the prediction that they would compete effectively for binding to the uridine-rich sequences often found in the polypyrimidine stretches near the 3' end of introns and downstream of the cleavage site for polyadenylation in pre-mRNAs. Moreover, evidence that suggested a role for hnRNP proteins in pre-mRNA splicing has been described (Choi et al. 1986; Sierakowska et al. 1986). Experiments using an in vitro RNA-binding assay demonstrated that three hnRNP proteins, A1, C, and D (Piñol Roma et al. 1988), show intrinsic binding specificities on authentic pre-mRNAs (Swanson and Dreyfuss 1988b). These proteins bind specifically to the 3' end of introns within a region containing the conserved polypyrimidine stretch between the branch site and the 3' splice site. The hnRNP C proteins also were found to cross-link by UV light preferentially to uridine-rich sequences downstream of the polyadenylation cleavage site on two different viral pre-mRNAs (Wilusz et al. 1988; Moore et al. 1988); this binding was sensitive to mutations that interfered with aspects of the in vitro cleavage–polyadenylation reaction (Wilusz et al. 1988).

In sum, contrary to an earlier view that hnRNP proteins coat pre-mRNA in a non-sequence-specific manner (Pullman and Martin 1983; Thomas et al. 1983; Wilk et al. 1983; Conway et al. 1988), these experiments demonstrate that several RNP-CS-containing hnRNP proteins do have RNA-binding specificities and can bind to sites on pre-mRNAs that are critical for the formation of mature and functional mRNA. Since the A1, C, and D proteins are confined to the nucleus, and appear to have the highest affinity for pre-mRNA sequences that are removed during the formation of mRNA, it is possible that these pre-mRNA segments constitute nuclear retention sequences (Swanson and Dreyfuss 1988a,b; Wilusz et al. 1988).

Developmental loci in Drosophila encode RNP-CS-type proteins
Several genes that appear to have critical roles in establishing certain developmental pathways in Drosophila and maize recently have been cloned molecularly and their sequences have predicted protein products that contain RNP-CS-type RNA-binding domains (Amrein et al. 1988; Bell et al. 1988; Gómez et al. 1988; Robinow et al. 1988; Goralski et al. 1989). In maize, the plant hormone abscisic acid induces in the embryo an RNP-CS containing protein whose function is unknown (Gómez et al. 1988; Mortenson and Dreyfuss 1989). The Drosophila genes, embryonic lethal abnormal visual system (elav), sex-lethal (Sxl), and transformer-2 (tra-2), are critical in the early determination, as well as in the maintenance, of two terminally differentiated states, the neuron and the somatic sexual state. Although it has not been shown yet that these gene products are RNA-binding proteins in vivo, considerable genetic and molecular data suggest that these loci exert their regulatory effects at the post-transcriptional level. Indeed, it appears that the entire sex-determination pathway in Drosophila is regulated at the level of sex-specific splicing (Boggs et al. 1987; Baker and Wolfner 1988; Nagoshi et al. 1988). Sxl positively regulates the activity of the feminizing gene, transformer (tra); the protein product of the Sxl locus is necessary to produce a correctly spliced, functional tra mRNA in females (Cline 1979; McKeown et al. 1988, Nagoshi et al. 1988). Although Drosophila males and females express Sxl transcripts, only females encode a mRNA capable of producing a full-length, functional polypeptide; male transcripts, on the other hand, harbor an exon with an in-frame translation termination codon. Bell et al. (1988) propose that Sxl transcripts are most likely subject to an autoregulatory, alternative splicing mechanism that maintains the appropriate sexually differentiated state, producing functional Sxl protein and correctly spliced tra mRNA in females only.

Role in alternative splicing
Considering that RNP-CS-containing proteins such as Sxl, tra-2, elav, and the abscisic acid-induced maize protein are most probably RNA-binding proteins, they could have any one of many possible functions in RNA metabolism. However, as the foregoing discussion suggests, particularly for Sxl and tra-2, a role in alternative splicing is likely and interesting models on how this might be accomplished by such proteins can be envi-
RNA-binding proteins


tioned. The occurrence of multiple, nonidentical RNA-binding domains in a single protein suggests that, like any macromolecule that has multiple ligand binding sites, RNP-CS-type RNA-binding proteins may be able to bind to multiple segments of RNA, with each domain possibly binding independently of the others. Together with the finding that RNP-CS-containing proteins can have RNA binding specificity, it appears that such proteins have the potential to bring into proximity segments of RNA that are otherwise far apart on the same RNA molecule or even on different RNA molecules. In this context, the binding preference of some hnRNP proteins for sites on the pre-mRNA that are important for pre-mRNA processing—the 3' splice site and sequences downstream of the polyadenylation site—is particularly tantalizing. An extension of these findings is the possibility that such proteins may play a role in pre-mRNA splicing, and if their modular structure contains the appropriate combination of binding specificities, also in alternative splicing. RNP-CS-type proteins potentially may affect alternative splicing by negative regulation as well. In this scheme, a protein with even a single RNA-binding domain could compete with other nuclear factors for a specific set of 3' splice sites, and it could contain an auxiliary domain(s) that prevents, rather than promotes, splicing complex formation at that site.

Post-transcriptional control mechanisms, including a myriad alternative splicing events, appear to operate in a growing number of cellular processes and pathways (for review, see Kozak 1988). As the above discussion suggests, RNP-CS-type proteins are excellent candidates for a novel class of regulatory proteins affecting these mechanisms. To date, the best characterized examples of alternative splicing pathways that have major developmental consequences are in Drosophila, among them Sxl and tra-2. It is tempting to speculate that the protein product of Sxl, an RNP-CS-type protein with multiple RNA-binding domains, effects sex-specific alternative splicing by binding to different sets of sequences on an individual pre-mRNA, consequently promoting or repressing the selection of splice sites. By being expressed appropriately and by using the potential for sequence preference within its multiple, independent RNA-binding domains, a regulator protein such as the Sxl gene product could influence the production of novel mRNAs significantly.

Similarity to transcription factors

Finally, the emerging picture of the structure and function of RNP-CS-type RNA-binding proteins bears an intriguing similarity to some characteristics of transcription factors. The binding of RNP-CS-type proteins to RNA may not require recognition of a strict consensus binding site because they bind to the 3' end of introns and to sequences downstream of the polyadenylation signal that vary in nucleotide sequence. The binding of OBP100, the enhancer-octamer DNA-binding protein, to degenerate consensus binding sites is reminiscent of this situation [Baumruker et al. 1988]. In addition, one consequence of the hypothetical role of RNP-CS-type proteins in alternative splicing is that differential sequence binding by such proteins on a RNA molecule could lead to a looping out of RNA between these binding sites. This would be similar to the ability of a well-characterized specific DNA-binding protein, phage λ repressor, to bind cooperatively at distal, nonadjacent sites upstream of operator DNA in the λ genome, with the result that DNA in between is looped out. The inhibition of transcriptional activity that follows recently was shown to be a direct consequence of distal repressor—repressor interaction in vitro and in vivo [Hochschild and Ptashne 1988]. Finally, the auxiliary domains of several of the RNP-CS-type RNA-binding proteins are similar in amino acid character to some activation domains of eukaryotic transcription factors. For example, there are similarities in amino acid character between the highly acidic domains of the hnRNP C proteins and the transcription factor GCN 4 [Hope and Struhl 1986], the glutamine-rich amino-terminal domain of elav and SP1 [Courey and Tjian 1988], and the proline-rich domains of the PABP and CTF/NFI [Bohmann et al. 1987].

If the proteins of the Drosophila elav and Sxl loci are shown, as expected, to be bona fide RNA-binding proteins, a new and experimentally tractable function of RNP-CS-type proteins will have been uncovered. It remains an exciting possibility that the repertoire of developmental regulators now has to be expanded to include RNA-binding proteins.

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