RESEARCH COMMUNICATION

Ubiquitous expression and embryonic requirement for RNA polymerase II coactivator subunit *Srb7* in mice

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Mammalian RNA polymerase II complexes and coactivators containing homologs of yeast *Srb/Med* proteins have been isolated recently from tissue culture cells. The yeast *Srb/Med* complex is involved in global gene expression and is essential, but it is not yet known if its mammalian counterparts are broadly expressed in tissues or if they are essential. We have isolated the murine gene encoding *Srb7*, an *Srb/Med* complex protein whose sequence and function is highly conserved between yeast and humans. The mouse *Srb7* gene is single copy, and Northern analysis showed that it is expressed in all tissues examined. Disruption of the gene in embryonic stem cells revealed that it is essential for cell viability and murine embryonic development. These results, together with evidence that murine *Srb7* is associated exclusively with high molecular weight forms of RNA polymerase II in extracts, suggest that *Srb7*-containing polymerase complexes occur in most tissues and have essential roles in expression of protein coding genes.

Transcription initiation in eukaryotes involves the recruitment of RNA polymerase II and associated factors to promoters by gene-specific activators (Orphanides et al. 1996; Roeder 1996; Ptashne and Gann 1997; Hampsey 1998; Myer and Young 1998). Attempts to reconstitute transcription in vitro led to the identification of one set of initiation factors that are associated with RNA polymerase II at promoters, and these were called basal or general transcription factors (GTFs). The GTFs include TFIID, TFIIB, TFIIE, TFIIIF, and TFIIH. Genetic and biochemical studies have identified additional factors that are necessary for appropriate regulated expression in vivo or for reconstitution of more physiologically relevant activities in vitro, among which are the yeast *Srb/Med* complex.

We and others have proposed that the form of RNA

polymerase II that is generally recruited to promoters by transcriptional activators in yeast cells contains an *Srb/Med* complex (for review, see Koleske and Young 1995; Ptashne and Gann 1997; Hampsey 1998; Myer and Young 1998). The genes encoding the yeast *Srb* proteins were discovered through a genetic screen designed to identify components of the transcription apparatus, which are involved in the response to transcriptional regulators (Nonet and Young 1989). Attempts to purify *Srb* proteins led to the isolation of a large complex containing core RNA polymerase II, a subset of the general transcription factors, and various regulatory proteins (Thompson et al. 1993; Koleske and Young 1994). This holoenzyme complex had the capacity to initiate transcription and respond to activators when supplemented with additional purified general transcription factors in vitro. A subcomplex that dissociated from the holoenzyme with anti-CTD antibodies was found to reconstitute the response to activators in vitro; this subcomplex contains *Srb* and additional proteins called *Meds* (Kim et al. 1994). The response to activators is critical because in vitro systems reconstituted with yeast GTFs and polymerase alone were not activator-responsive (Flanagan et al. 1991). Two of the yeast *Srb* proteins (*Srb4* and *Srb6*) were found to be required for transcription of most protein coding genes in vivo (Thompson and Young 1995; Holstege et al. 1998). Because these *Srb* proteins are found tightly associated with the holoenzyme, it follows that the *Srb*-containing holoenzyme is the form of RNA polymerase II that is recruited to most promoters in yeast cells.

Transcription-competent RNA polymerase II complexes containing *Srb* homologs were identified in mammalian cells soon after the discovery of the yeast holoenzyme (Ossipow et al. 1995, 1999; Chao et al. 1996; Maldonado et al. 1996; Pan et al. 1997; Cho et al. 1998; Neish et al. 1998). The precise composition of these complexes was not established, due in part to technical difficulties associated with purifying intact megadalton size multiprotein complexes. However, mammalian *Srb7* was found to be a component of all complexes assayed by Western blot and was shown to be tightly associated with high molecular weight forms of RNA polymerase II (Ossipow et al. 1999). More recently, investigators have described more highly defined protein complexes that were isolated using assays for cofactor function and that contain homologs of the yeast *Srb/Med* proteins (Jiang et al. 1998; Gu et al. 1999; Naar et al. 1999; Rachez et al. 1999; Ryu et al. 1999). These purified complexes lack RNA polymerase II but associate with the enzyme where they function as coactivators and corepressors (Hampsey and Reinberg 1999).

These recent developments suggest that *Srb/Med*-containing complexes have diverse and important roles in regulation of transcription initiation in mammalian cells. It would be interesting to determine whether *Srb/Med* genes are generally expressed in mammalian tissues and whether they are essential for viability. To address

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these issues, we have isolated the murine *Srb7* gene and used it to determine the range of mammalian tissues that express this gene and to ascertain whether it is essential for cell viability and development.

Results and Discussion

Murine *Srb7* genomic and cDNA clones were isolated from libraries by using a human *SRB7* cDNA clone (Chao et al. 1996) as probe. Among the *Srb*/mediator proteins shared by yeast and mammalian holoenzymes, *Srb7* was selected for investigation because its sequence and function is highly conserved between yeast and humans (Chao et al. 1996), it has been observed in all mammalian RNA polymerase II complexes examined thus far (Chao et al. 1996; Maldonado et al. 1996; Scully et al. 1997; Neish et al. 1998; Ossipow et al. 1999), and *Srb7* is associated exclusively with the high molecular weight form of RNA polymerase II in murine cell extracts (Ossipow et al. 1999). Southern analysis of genomic DNA indicated that the *Srb7* gene is single copy (data not shown). The cDNA clone and a portion of the genomic clone were sequenced, and the predicted mouse *Srb7* protein sequence is shown in Figure 1. The 144-amino-acid mouse and human *Srb7* proteins are identical except for a single amino acid difference at residue 36. Orthologous sequences were also identified in *Caenorhabditis elegans* and *Schizosaccharomyces pombe* (Fig. 1). Sequence homology among the *Srb7* proteins was highest in the amino terminus, which is the portion of human *Srb7* which can functionally substitute for the corresponding yeast sequence (Chao et al. 1996).

The presence of *Srb7* mRNA in various tissues was

H.sapiens	1	MADR	L	TQLQDAVNS	LADQFCNA	I	IGVL	-	-	-	QQCG	30
M.musculus	1	MADR	L	TQLQDAVNS	LADQFCNA	I	IGVL	-	-	-	QQCG	30
C.elegans	1	MADR	M	TQLQDMINEMAG	MTNA	I	IGVL	-	-	-	QATA	30
S.pombe	1	MACR	C	TQLQDITIDEVA	TQFYSS	I	IHYLS	SSH	H	F		33
S.cerevisiae	1	MTDR	L	TQLQCLDQMTE	QFCAT	L	N	YTD	K	NH	GFE	33
H.sapiens	31	P	P	ASFNN	IQTAINKDQ	PAN	PTEEYA	QAL	FAAL	I		62
M.musculus	31	P	P	ASFNN	IQTAINKDQ	PAN	PTEEYA	QAL	FAAL	I		62
C.elegans	31	P	P	CFEFGA	ISGELEDEPN	-	-	-	-	CAIF	AAG	55
S.pombe	34	PL	T	PGQEKVSDSKVN	PISAEEL	QFA	QRDL	A	K	D		65
S.cerevisiae	34	RL	T	VNEPQMSDKHATVV	PEEF	FSNT	IDEL	S	T	D		66
H.sapiens	63	ARTAKD	I	DVIL	ISLPS	EE	-	STAAL	QAAS	LYKLE		94
M.musculus	63	ARTAKD	I	DVIL	ISLPS	EE	-	STAAL	QAAS	LYKLE		94
C.elegans	56	AKAAKN	IE	IL	IS	EP	EAGN	QEA	VEE	KMIKND		88
S.pombe	66	VT	KFMQ	ITD	INQL	PGIS	TAPKH	QLE	IKK	KLQ		97
S.cerevisiae	67	IL	KTRQ	IN	KL	ISL	PGVD	VSAE	QL	RKID	MLQ	98
H.sapiens	95	EENH	E	AATCL	EDVVYRGDM	L	EKI	QSAL	ADIAQ			127
M.musculus	95	EENH	E	AATCL	EDVVYRGDM	L	EKI	QSAL	ADIAQ			127
C.elegans	89	EKQRE	K	VNEL	VGLVGD	SN	RL	VGV	VQKKL	AEISK		121
S.pombe	98	NS	IEE	KQLER	KSLESE	NE	DL	KLQ	LAKRI	ETFG		130
S.cerevisiae	99	KKLV	E	VEDE	KIEAT	KKKEK	ER	L	LRH	VD	SLIEDFVD	131
H.sapiens	128	S	QLK	TRSGTHS	QS	SL	PD	S				144
M.musculus	128	S	QLK	TRSGTHS	QS	SL	PD	S				144
C.elegans	122	T	QMSSRP	-	-	SE	-	-	-	-	-	130
S.pombe	131	L	SCVLFQ	P	-	-	-	-	-	-	-	138
S.cerevisiae	132	G	I	ANS	K	K	S	T	-	-	-	140

Figure 1. Alignment of sequences predicted for mouse, human, worm, and yeast *Srb7* proteins. Human and *S. cerevisiae* *Srb7* protein sequences are from Chao et al. (1996) and Myers et al. (1998), respectively. The putative *S. pombe* and *C. elegans* homologs are derived from GenBank accession nos. 4007762 and Z81475 (gene C24H11.6, aligned sequence starting at conserved methionine), respectively. The asterisk (*) indicates the single amino acid difference between human and mouse *Srb7*. Boxed residues are identical; shaded residues are similar as determined by the SeqVu program (Riek et al. 1995).

Figure 2. Expression of *Srb7*. The full-length *Srb7* cDNA clone was hybridized to a multiple tissue Northern blot from Clontech.

investigated by probing a multi-tissue Northern blot with radiolabeled *Srb7* cDNA (Fig. 2). A single band of the predicted size was observed in samples from each of the tissues examined, including heart, brain, spleen, lung, liver, muscle, kidney, and testes. We conclude that the *Srb7* gene is expressed in most murine tissues.

The requirement for *Srb*/Med components in vivo has not been established in higher eukaryotes. We disrupted the *Srb7* gene in embryonic stem (ES) cells to explore the requirement for its function in the mouse. A genomic DNA clone was used to create a targeting construct that deletes the second and third exons of the *Srb7* gene (Fig. 3A), eliminating sequence coding for residues 16–87 of the 144-amino-acid reading frame. ES cells were transfected with the targeting vector and drug-resistant colonies were screened for homologous integration, which occurred (Fig. 3B, left) in ~5% of the clones. These cells showed no obvious growth phenotype. To determine if a homozygous deletion of the *Srb7* gene could be achieved, heterozygous ES cell clones were retargeted with a vector containing a different drug resistance marker. The mutated allele was replaced (Fig. 3B, right) at a frequency similar to that seen in the initial targeting experiments (9/74), whereas no clones were found to be homozygously disrupted (0/74). This is consistent with inviability of homozygous ES cells and suggests that the *Srb7* gene is essential for cell viability.

To further explore the effects of *Srb7* disruption, heterozygous ES cells were used to derive mice mutant for *Srb7*. Of six chimeras derived from three different ES clones, one animal, which showed 40% chimerism, gave germ-line transmission. Heterozygous animals showed no overt phenotype. In contrast, no homozygous animals were born from heterozygous intercrosses, suggesting that the absence of *SRB7* is lethal.

Next we examined the developmental stage at which homozygous embryos die. As embryonic transcription begins soon after cleavage commences, it would be expected that maternal stores of the *Srb7* protein would allow the zygote to reach at least this stage. To test this, embryos of different developmental stages were genotyped by PCR (Fig. 4). Homozygous embryos were found

Figure 3. (A) Targeted disruption of *Srb7*. An 8.5-kb *Bam*HI–*Eco*RI (E and B) genomic fragment in pBS II KS[−] was the basis for the targeting construct. Exons II and III were replaced by a PGK–neo or PGK–hyg cassette. Exons are numbered I–IV; construct pMT204 is the same as pMT203, except the drug resistance marker was replaced with a PGK–hyg cassette. (B, left) *Bam*HI-digested genomic DNA from G418-resistant ES cell clones (transformed with pMT203) were genotyped by Southern hybridization with the probe indicated in A. (Right) Clones targeted with the neo-containing vector (203-targeted) were retargeted with the hyg vector (204-targeted). Hygromycin B-resistant clones showed replacement of the *neo* allele but not disruption of the remaining wild-type allele.

in Mendelian ratios ($n = 26$, $\chi^2 P < 0.05$) until the blastocyst stage. Culture of blastocysts yielded no outgrowth of homozygous cells, suggesting developmental arrest at this stage (data not shown). Analysis of later embryos (7.5 and 8.5 dpc) showed numerous empty decidua and no homozygous embryos ($n = 25$, significant departure from Mendelian ratios, $\chi^2 P < 0.01$). We conclude that the *Srb7* gene product is essential for both embryonic development and cell viability. The survival of embryos to the blastocyst stage is likely due to maternal stores of the protein. By comparison, homozygous disruptions of *Rad51* die immediately after the beginning cleavage (Tsuzuki et al. 1996), whereas cyclin A2^{−/−} embryos sur-

vive to the early postimplantation stage though the deletion is cell lethal (Murphy et al. 1997). Whereas *Rad51*, cyclinA2, and *Srb7* disruptions are all apparently cell lethal, they have disparate developmental phenotypes, the latter being temporally intermediate. Compared to disruptions of other transcription factors, *Srb7*^{−/−} embryos show earlier lethality (−4–6 dpc) than those deficient for the abundant transcription factor *Sp1* (11 dpc) (Marin et al. 1997), the transcriptional coactivator/histone acetylase p300 (9 dpc) (Yao et al. 1998), or the chromatin remodeling factor *BRM* (nonlethal proliferation defects) (Reyes et al. 1998).

Recent studies have identified mammalian RNA polymerase II coactivator complexes containing *Srb*/Med proteins, but it has not been clear whether these complexes have general or essential roles in gene expression, as is the case in yeast cells (Holstege et al. 1998). The results described here reveal that the gene encoding *Srb7* is expressed in most mouse tissues and is essential for viability, suggesting that *Srb7*-containing polymerase complexes exist in most tissues and play essential roles in expression of protein coding genes.

Materials and methods

Isolation of clones and expression patterns

The human cDNA clone was used to probe a mouse brain cDNA library [provided by Tony Koleske (Yale University, New Haven, CT) and David Baltimore (California Institute of Technology, Pasadena, CA)] by hybridization, and positive clones were sequenced. The cDNA was then used to probe a 129 SVI BAC genomic library (Genome Systems) to isolate overlapping genomic clones. A 10-kb *Bam*HI fragment was found to contain the entire coding sequence and was subcloned into pBluescript II (PM429). This segment was used for subsequent constructions. Exons were identified by restriction mapping and partial sequencing of PM429.

The cDNA clone was hybridized to a mouse multitissue Northern blot (Clontech).

Targeting

Either a PGK–neo or a PGK–hyg selection cassette was inserted between the *Avr*II and *Hpa*I sites of an 8.5-kb genomic subclone, resulting in the deletion of the second and third exons of the gene [pMT203 and pMT204,

Figure 4. (A) Schematic depicting PCR genotyping assay. The number 104 denotes primer oMT104, common to both alleles; 103 denotes oMT103, specific to the wild-type allele; 106 denotes oMT106, specific to the mutant allele. (B) PCR genotyping of 3.5 dpc embryos (blastocysts or late morulae) derived from heterozygous crosses. Embryos were digested individually with proteinase K and subjected to PCR as detailed in Materials and Methods. Control reaction contained media alone. Wild-type, heterozygous, and homozygous mutant embryos were found in Mendelian ratios.

respectively). Linearized vectors were electroporated into J1 ES cells (Li et al. 1992), and drug-resistant colonies were screened by PCR and Southern hybridization of *Bam*HI-digested DNA with an external probe (pMT205). Homologously targeted clones were shown to be single copy integrants by probing with an internal probe. Retargeting experiments were analyzed by Southern hybridization. In these experiments the individual alleles could be distinguished by restriction polymorphisms. This allowed the recognition of targeting events that replaced one mutant allele with another. Heterozygous clones were injected into BALB/c blastocysts, and resulting animals chimeric by coat color (20%–90%) were mated with BALB/c and 129Sv¹ mice to generate mixed-background and inbred lines carrying the mutation. To genotype embryos, heterozygotes were mated, embryos were collected and developmental stage noted, and were placed in 0.2-ml PCR tubes with 5 μ l of proteinase K buffer (1 $\times$ Promega Mg-free PCR buffer + 200 μ g/ml proteinase K). After 1 hr digestion at 50°C, the proteinase K was inactivated by heating to 95°C for 10 min, a PCR master mix was added, and the samples were amplified and analyzed by gel electrophoresis. Southern hybridization with an internal oligonucleotide was used to confirm identity of PCR products. The following primers were used: oMT103, GGGAGGCATTTGAGGCTTATTGATTTTG (wild-type allele); oMT104, CACCAC-CACCAACCCAGCTGCTCTA (common); oMT106, CTTAATCGCCTT-GCACCACATCCC (mutant allele); and oMT108, CCTCTGGAAGAT-TGTGCTGT (internal hybridization oligonucleotide).

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