DPY-30, a nuclear protein essential early in embryogenesis for *Caenorhabditis elegans* dosage compensation

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SUMMARY

DPY-30 is an essential component of the *C. elegans* dosage compensation machinery that reduces X chromosome transcript levels in hermaphrodites (XX). DPY-30 is required for the sex-specific association of DPY-27 (a chromosome condensation protein homolog) with the hermaphrodite X chromosomes. Loss of dp y-30 activity results in XX-specific lethality. We demonstrate that dp y-30 encodes a novel nuclear protein of 123 amino acids that is present in both hermaphrodites and males (XO) throughout development. DPY-30 itself is not associated with the X chromosomes, nor is its pattern of expression perturbed by mutations in the gene hierarchy that controls dosage compensation. Therefore, DPY-30 is a ubiquitous factor that is likely to promote the hermaphrodite-specific association of DPY-27 with X by affecting the activity of a sex-specific dosage compensation gene. In XO animals, DPY-30 is required for developmental processes other than dosage compensation: coordinated movement, normal body size, correct tail morphology and mating behavior. We demonstrate that rescue of both the XX-specific lethality and the XO-specific morphological defects caused by dp y-30 mutations can be achieved by inducing dp y-30 transcripts either in the mother or in the embryo through the end of gastrulation. dp y-30 appears to be cotranscribed in an operon with a novel RNA-binding protein.

Key words: dosage compensation, X chromosome, *Caenorhabditis elegans*, gene expression, RNA-binding protein

INTRODUCTION

Dosage compensation is a chromosome-wide regulatory process that controls the expression of numerous genes related solely by their linkage to the same sex chromosome. The need for dosage compensation arises in organisms whose primary sex-determining mechanisms cause males and females to differ in their relative dose of X chromosomes. Dosage compensation mechanisms equalize expression of X-linked genes between the sexes and prevent the sex-specific lethality that would otherwise result from the two-fold difference in X-linked gene dose. The mechanisms used to achieve dosage compensation are diverse. In mammals, dosage compensation is accomplished by random inactivation of one of the two female X chromosomes, thereby reducing the effective X chromosome dose to that of males (XY) (reviewed in Migeon, 1994). In *Drosophila*, the dosage compensation machinery acts to transcribe the single X chromosome of males (XY) at twice the rate as each of the two X chromosomes in females (XX) (reviewed in Baker et al., 1994). In *C. elegans*, XX hermaphrodites reduce the transcript levels produced by each X chromosome to achieve the same levels produced by the single X of males (XO) (reviewed in Hsu and Meyer, 1993).

Recent molecular characterization of genes essential for dosage compensation in mammals, flies and worms has greatly advanced our understanding of these unique global regulatory mechanisms. Studies in mammals have identified a large, non-coding transcript (*Xist*) from within the cis-acting X chromosome inactivation center that is specific to the inactive X and may play a role in initiating X chromosome inactivation (Brockdorff et al., 1992; Brown et al., 1992; Brown and Willard, 1994; Kay et al., 1993). In *Drosophila*, the proteins essential for elevated transcription of the male X (MLE, a putative RNA helicase, MSL-1 and MSL-3, two novel proteins, and MSL-2, a ring finger protein) are sex-specifically localized to the male X, a finding consistent with their direct involvement in X chromosome hypertranscription (Kuroda et al., 1991; Palmer et al., 1993; Gorman et al., 1995; Kelley et al., 1995; Zhou et al., 1995). The association of these proteins with X requires the functional products of all the *msl* and *mle* genes, suggesting that they form a multimeric complex. In addition, an acetylated isoform of histone H4 colocalizes with the MSL and MLE proteins on X in a manner dependent on their activities (Turner et al., 1992; Bone et al., 1994), suggesting that the male dosage compensation apparatus modulates gene expression through modification of X chromatin structure. In *C. elegans*, insight into how X chromosome expression is reduced in hermaphrodites has been gained through analysis of the hermaphrodite-specific dp y-27 dosage compensation gene (Chuang et al., 1994). Immunolocalization studies of DPY-27 showed that it is expressed in the nuclei of both sexes, but is sex-specifically localized to only the hermaphrodite X chromosomes, consistent with its role in down regulating X chromosome expression. DPY-27 belongs...
expression, but they do cause XO animals to have numerous morphological and behavioral abnormalities, including a delay in development, a small body size, an inability to mate and a defect in male tail structures. The pleiotropic phenotypes caused by *dpy-30* mutations suggest that in addition to acting in dosage compensation, *dpy-30* plays a more general role in the development of both XX and XO animals.

We show that *dpy-30* encodes a novel nuclear protein ubiquitously expressed throughout the development of both XX and XO animals. In contrast to *DPY-27*, *DYP-30* is not associated with the X chromosome, nor is its pattern of expression perturbed by mutations that disrupt dosage compensation, or by changes in the X:A ratio. Therefore, *DYP-30* is likely to be required for the activity of sex-specific dosage compensation factors that down regulate X-linked gene expression in XX animals. Our studies also indicate a requirement for *DYP-30* during early embryogenesis in XX animals, consistent with the need for an active dosage compensation process early in development. XO animals have a similar temporal requirement for *DYP-30*. Lastly, *dpy-30* appears to be cotranscribed in an operon with a novel RNA-binding protein.

**MATERIALS AND METHODS**

**C. elegans cultures**

To obtain large populations of nematodes of all developmental stages for RNA extractions, animals were grown in liquid culture and isolated as described in Klein and Meyer (1993). Adult XO males were isolated by the Nytex screening procedure described by Meyer and Casson (1986).

**Germline transformation**

Germline transformation rescue experiments were carried out as described in Mello et al. (1991) by microinjecting *dpy-30(y130)* young adults with a solution containing both test DNA (10 µg/ml) and marker DNA (plasmid pRF4, 100 µg/ml), which carries the dominant rol-6 (su1006) marker and confers a roller (Rol) phenotype. Healthy *dpy-30(y130)* homozygous animals were obtained for injection as the non-Unc progeny of heterozygous *dpy-30(y130) / nT1(n754dm)* mothers. Cosmids and subclones were assayed for their ability to rescue the XX-specific lethality by examining broods of injected *dpy-30* XX animals for viable F1 roller progeny. If F1 rescue was obtained, rescued animals were picked to individual plates to establish stably transmitting lines carrying the rescuing construct. Constructs were eliminated from consideration if injections of 50 to 100 hermaphrodites failed to produce any viable progeny. Initial transformation experiments with the cosmids C02C8, C08F12, C09F3, C27A7, C35G11, C51B3 and F29D2 (from A. Coulson and J. Sulston) demonstrated that C09F3, located 50 kb to the left of the myo-3-cosmid C02B11, rescued the *dpy-30* XX transgenic animals. Stable arrays made from subclones of C09F3 (Fig. 1) were used to delineate the minimal genomic region required for rescue of the *dpy-30* mutant phenotypes. For each rescuing construct, at least two independent transmitting lines were scored for rescue of the *dpy-30* XX-specific phenotypes. The smallest rescuing clone, pDH38, also rescues the XX-specific lethality and the XO-specific morphological defects caused by the *dpy-30(y228)* null allele.

**Isolation of RNA and northern hybridization experiments**

Total RNA and poly(A)* RNA were made as described by Klein and Meyer (1993) and Hsu and Meyer (1994). Northern analysis was performed as described by Meyer and Casson (1986). The *dpy-30*-specific probe was the 1.3 kb *Hincl* to *Sst* fragment from pDH13.
that contains the entire dpy-30 rescuing region. After probing with dpy-30, blots were rehybridized with a probe for the act-1 gene (Krause et al., 1989).

Sequence analysis of dpy-30 and rnp-1
Sequence analysis of the 4 kb genomic DNA that contains the dpy-30 rescuing region, was performed according to Sambrook et al. (1989), using pDH13 and its reverse orientation clone pDH14. Additional genomic sequence 3' to dpy-30 was obtained from pDH8, a clone containing the entire rnp-1 genomic region (see Fig. 1). Nucleotide number 1 in the genomic sequence listed in Fig. 3 is the first base of the Xba I site at the beginning of the insert of pDH38 (Fig. 1C).

Three independent dpy-30 cDNA clones were obtained by probing N2 mixed-stage libraries (S. Kim and S. Lichtsteiner) and a him-8 embryonic library (L. Miller) with either a 32P-labeled 1.6 kb HindIII fragment from pDH12 or a 32P-labeled 1.3 kb HindII Styl fragment from pDH13. pDH42, the largest cDNA clone, begins at nucleotide 672 and ends at nucleotide 1323, with a poly(A)+ tail. pDH19 begins at nucleotide 826 and ends at nucleotide 1322, with a poly(A)+ tail. pDH20, begins at nucleotide 665 and continues to the EcoRI site at nucleotide 1057. The 5' end of pDH20 contains a 12 nucleotide sequence (5'CCCAAGTGTGAG 3') identical to the SL1 transspliced leader sequence found at the 5' end of many C. elegans mRNAs (Blumenthal, 1995). The trans-splicing pattern of dpy-30 was confirmed by PCR amplification of cDNA (Klein and Meyer, 1993) using oligonucleotides complementary to SL1 or SL2 and an oligonucleotide internal to dpy-30. DH-6 (5'CTGTGTTAGCTCCATTG-GCAG 3'). Using SL1 and DH-6, this amplification reaction produced an expected product of approximately 200 bp. No product was obtained using SL2 and DH-6.

The rnp-1 cDNA pDH18 was isolated fortuitously because the 1.6 kb HindII genomic probe used to isolate dpy-30 cDNAs also contained the first two exons of the rnp-1 transcript. This cDNA begins at nucleotide 1421 with the ATG initiation codon and ends with a poly(A)+ tail at nucleotide 3511 in the extended genomic sequence. The trans-splicing pattern of rnp-1 was determined by PCR amplification of cDNA using oligonucleotides complementary to SL1 or SL2 and an oligonucleotide internal to rnp-1, DH-9 (5'GCCGTGCTGCCAATTGGTGA 3'). The amplification reaction produced a product of approximately 300 bp using SL2 and DH-9. DNA sequence analysis of the cloned product showed that SL2 is spliced directly to the initiator ATG at nucleotide 1421. No product was obtained using SL1 and DH-9. Genomic DNA clones were sequenced on both strands; cDNA clones were sequenced on a single strand.

Sequence analysis of dpy-30 mutations
dpy-30 DNA was obtained from four homozygous dpy-30(y130 or y228) adult animals using the PCR method described by Klein and Meyer (1993) and the oligonucleotides DH-1 (5'CGAC- GAACAGTTGATGGCCATC 3') and DH-2 (5'GTCCTAGAGCAGCGTGTAACGTAC 3') to amplify dpy-30 products. These conditions yielded a single 1.4 kb product, consistent with the predicted size of the genomic fragment that should be obtained with DH-1 and DH-2. The DNA sequence of at least two independent clones derived from each amplification reaction was determined.

Construction of insertion mutations in the dpy-30 coding region
The insertion mutation within pDH43 was made by cutting pDH38 with Bcl I, filling in the 4 nucleotide overhang with E. coli DNA polymerase I large fragment (Klenow) (Sambrook et al., 1989) and recircularizing the linear blunt-ended DNA. The presence of the 4 base pair insertion at the Bcl I site was confirmed by DNA sequence analysis. pDH9, which contains a frameshift mutation at the EcoRI site in dpy-30, was made and confirmed in similar fashion.

Sequence similarity searches
The initial analysis of the dpy-30 and rnp-1 nucleotide and protein sequence was performed using DNA Strider (Marck, 1988). Similarity searches for the dpy-30 predicted protein were performed using either the BLAST algorithm (Altschul et al., 1990) or the FASTA algorithm (Devereux et al., 1984).

Recombinant DPY-30 protein, DPY-30 antibodies and immunocytochemistry
Expression of full-length recombinant DPY-30 protein and generation and analysis of DPY-30 antibodies were performed as in Chuan et al. (1994). For in situ antibody staining, embryos, larvae and adults were prepared as described in Chuan et al. (1994) from the following strains: wild-type XX, sdc-1(m485) XX, dpy-21(e428) XX, sdc-3(y129) XX, dpy-27(y167) XX, dpy-28(y1ts) XX, dpy-28(s939); him-5(e1490); xol-1(y9) XO, dpy-2(s202ts) XX, her-1(lv1y101); xol-1(y9) sdc-2(y82) unc-9(e101) XO, her-1(lv1y101); xol-1(y9) sdc-2(y74) unc-9(e101) XO and him-8(e1489) yh2[xol-l::lacZ, rol-6(su1006)]. To obtain XX embryos homozygous for either dpy-30(y228ts) or dpy-28(y1ts), homozygous mutant XX animals were first grown for several generations at the partially permissive temperature (15°C) and then shifted to the non-permissive temperature (20°C) for at least one generation. The gravid hermaphrodites were then treated with hypochlorite. The same regime was used for the sdc-2(y202ts) strain, except the temperature shift was from 20° to 25°C. dpy-27(y167) XX embryos were obtained as described in Chuan et al. (1994). The procedures for antibody staining, epifluorescence and confocal microscopy are as described in Chuan et al. (1994).

hsp16-41::dpy-30 fusion
To determine the time in development when dpy-30 transcription is essential for the viability of XX animals, dpy-30(y228) embryos or ooocytes carrying an integrated heat-inducible hsp16-41::dpy-30 fusion (yIs15) were heat shocked at different stages of their development (Rhind et al., 1995; Sulston et al., 1983). For heat shocks applied to embryos younger than comma stage, plates containing 30-40 gravid, non-Egl yIs15; dpy-30(y228) young adult hermaphrodites from dpy-30+ mothers were incubated at 20°C for 3 to 12 hours. These plates, which contained mixed-stage embryos, were then shifted from 20°C to 30°C for 30 minutes. Subsequently, comma-stage embryos were collected for 30 minute intervals to plates pre-cooled to 20°C. Animals collected by this method vary in age by approximately 45 minutes. For heat shocks applied after comma stage, comma-stage cohorts were collected and allowed to develop for defined intervals before applying the heat shocks. The time of heat shock relative to comma stage was defined as the time between the midpoint of the 30 minute heat-shock period and the midpoint of the 30 minute collection interval. Heat-shock times were subsequently expressed relative to first cleavage. 5 days after embryos were collected, the plates were scored for the presence of adult hermaphrodite progeny. For most time points, between 50 and 175 embryos were scored. Percent viability was calculated according to the formula: Viability = 100 (number of viable adult hermaphrodites)/(total number embryos). The baseline viability of dpy-30(y228) XX animals in the absence of heat shocks is approximately 12% (Hsu and Meyer, 1994). When survivors were present, they were also scored for the severity of the Dpy, Egl and Pvu dosage compensation phenotypes. The percent of the survivors that were wild type in length was calculated as follows: percent wild type = 100 (number of wild-type length survivors)/(total number of survivors).

To determine the time between 3 and 10 hours of embryonic development that transcription of dpy-30 is required to rescue the dpy-30 XO-specific morphological defects, groups of 20 yIs15; dpy-30(y228) him-5(e1490) young non-Egl adult hermaphrodites from dpy-30+ parents were placed on fresh plates pre-cooled to 20°C, allowed to lay embryos for 1 hour, and transferred to fresh pre-cooled plates. Cohorts...
collected this way were allowed to develop for defined periods of time and then were heat shocked as described above. Embryos synchronized by these brood collections vary in age by up to 3 hours. 3-5 days after brood collections, adult males were scored for rescue of the small (Sma), uncoordinated (Unc) and disrupted tail phenotypes caused by the dpy-30 mutation. The time of the heat shock was defined as the time between the midpoint of the hour long brood collection and the midpoint of the 30 minute heat shock. Heat-shock times were subsequently rescaled to reflect the time relative to first cleavage based on the observation that yIs15; dpy-30(y228) him-5(e1490) embryos are laid approximately 4 hours prior to comma stage. To induce dpy-30 transcription prior to 3 hours of embryonic development, non-Egl gravid yIs15; dpy-30(y228) him-5(e1490) hermaphrodites were shifted to 30°C for 30 minutes and their progeny subsequently collected at defined times and scored as described above. For most time points in both sets of experiments, between 22 and 53 males were scored. The percent wild-type males was calculated according to the formula: Percent wild type = 100 (number of wild-type males in a cohort)/(total number of males). In the absence of heat shocks, approximately 17% of the XO animals produced by yIs15; dpy-30(y228) him-5(e1490) hermaphrodites resemble wild-type XO males.

RESULTS

Molecular cloning of the dpy-30 dosage compensation gene

We cloned dpy-30 using germline transformation rescue experiments in conjunction with C. elegans genetic and physical map data. Genetic analysis had placed dpy-30 near the center of chromosome V, approximately 0.1 map units to the left of the previously cloned myosin heavy chain gene myo-3 (Miller et al., 1986) and to the right of the actin gene cluster act-1, 2, 3 (Krause et al., 1989), in a 600 kb region of the genome that is represented by a contiguous set of overlapping cosmids clones (Coulson et al., 1986; Hsu and Meyer, 1994) (Fig. 1A). Cosmids to the left of myo-3 were assayed for the presence of dpy-30(+) by testing their ability to rescue the dpy-30 XX-specific maternal-effect lethality in germline transformation experiments (Materials and Methods). These experiments identified a single cosm, C09F3, that rescues the XX-specific lethality caused by the dpy-30(y130) allele (Fig. 1B). Additional transformation experiments assaying subclones and deletions of C09F3 further delimited the dpy-30 rescuing region to a small, 921 bp interval contained within pDH38 (Fig. 1B,C).

The rescue obtained with all subclones of this region was qualitatively similar to that obtained using the entire C09F3 cosmid, suggesting that most, if not all, of the essential elements of the dpy-30 gene are contained within the 921 bp region. Subsequently, we determined that pDH38 also rescues animals homozygous for the dpy-30(y228) null mutation. In addition, pDH38 rescues the uncoordinated (Unc), small (Sma) and disrupted tail phenotypes associated with dpy-30 XO males (data not shown).

dpy-30 produces a single small transcript that is expressed throughout development

To determine the temporal pattern of dpy-30 mRNA expression in XX animals, we performed northern hybridization experi-

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*Fig. 1.* Genetic and physical location of dpy-30. (A) Genetic map of the center of chromosome V. dpy-30 lies in the 0.4 map unit interval between SMA-1 and myo-3 (Hsu and Meyer, 1994). The flanking gene myo-3 (bold lettering) and the act-1, 2, 3 gene cluster (bar below map) define a region of approximately 600 kilobases that must contain dpy-30. Chromosomal deficiencies yDf12 and ctDf1 are drawn below the map. (B) Rescue of dpy-30 mutant phenotypes with subclones of cosmid C09F3. The top line shows a partial restriction map of dpy-30 rescuing cosmid C09F3. Horizontal lines under the restriction map show the extents of various subclones of this genomic region. The ability of each subclone to rescue dpy-30 mutants is indicated by a plus and a black line (rescue) or a minus and a gray line (failure to rescue). The dashed vertical lines and the gray filled rectangle indicate a 4 kb region of genomic DNA encoding dpy-30 rescuing activity. Enzymes shown are: PvuI (PA), KpnI (K), BssHII (BS), PstI (PS), SalI (S), NdeI (N), HpaI (H). (C) Additional rescue experiments limit the dpy-30 rescuing region to approximately 1 kb. pDH38, a 1.3 kb genomic clone, rescues all dpy-30 phenotypes in XX and XO animals. The mature dpy-30 transcript is shown; rectangles represent exons, lines represent introns and shaded regions indicate coding regions. Enzymes shown are: XbaI (X), PvuI (PV), EagI (EA), ApaLI (A), EcoRV (RV), BclI (B), EcoRI (E), NdeI (N), ClaI (C), StyI (S), BspMI (BS), BglII (BG).
ments in which poly(A)+ RNAs prepared from synchronized populations of wild-type embryos, the four larval stages and adults were probed with genomic DNA spanning the dpy-30 locus. As shown in Fig. 2A, a single, relatively abundant 0.6 kb transcript that is expressed at approximately equal levels in all developmental stages was detected. This ubiquitous expression pattern is consistent with a role for dpy-30 in the initiation and maintenance of the XX-mode of dosage compensation.

To address whether any sex-specific regulation of dpy-30 is apparent at the transcript level, we also probed northern blots carrying either total RNA from purified populations of adult males or poly(A)+ RNA prepared from a population of him-8 embryos, consisting of approximately 37% XO embryos and 63% XX embryos (Hodgkin et al., 1979). The population of XX and XO embryos express the 0.6 kb dpy-30 transcript at a similar level to that seen in XX embryos (Fig. 2A). XX and XO adults also express similar dpy-30 transcript levels (Fig. 2B). The expression of dpy-30 in animals of both karyotypes is consistent with genetic evidence demonstrating that dpy-30 is required in both sexes (Hsu and Meyer, 1994). The absence of additional transcripts in XO animals indicates that the 0.6 kb dpy-30 transcript is likely to encode a single protein that is required in both sexes.

The dpy-30 gene encodes a small, novel protein

To determine the nucleotide sequence and splicing pattern of the dpy-30 transcript, we analyzed three independent dpy-30 cDNA clones and compared their DNA sequence to that of genomic DNA from the 4 kb rescuing clone pDH13 (Materials and methods). These partial, overlapping cDNA clones all correspond to the 0.6 kb transcript detected on northern blots. The 5’ end of the transcript was identified by virtue of one cDNA clone containing a short 12 nucleotide sequence 5’ CCCAAGTTTGAG 3’ that is absent from the genomic dpy-30 DNA but is identical to part of the SL1 trans-spliced leader sequence found at the 5’ end of many C. elegans mRNAs (Blumenthal, 1995). To confirm that the mature dpy-30 mRNA is trans-spliced, we performed RT-PCR using oligonucleotides for SL1 and a region internal to dpy-30 (Fig. 3). The reaction produced a single product that corresponds to the size predicted by the existing trans-spliced cDNA clone. A similar experiment using an oligonucleotide for the SL2 trans-spliced leader and the dpy-30 internal primer produced no product.

The mature dpy-30 transcript consists of two small exons forming a single small open reading frame encoding a protein of 123 amino acids (Fig. 3). A BLAST (Altschul et al., 1990) comparison of the dpy-30 protein sequence using the NCBI database revealed no significant similarities to known proteins or structural motifs, indicating that dpy-30 encodes a novel protein. Because of the small size of this predicted protein, we explored the possibility that dpy-30 might not encode a protein, but rather a regulatory or structural RNA that is required for the proper regulation of dosage compensation. To test this possibility, we engineered insertions of four nucleotides into the dpy-30 coding region after codon 3 (at a BclI site) or codon 99 (at an EcoRI site) (Fig. 3) and assayed their ability to rescue dpy-30 XX animals (see Materials and methods). These mutations, which change the reading frame and cause the new predicted proteins to be truncated after amino acids 19 or 103, prevent the altered genes from rescuing the dpy-30 XX-specific lethality, suggesting that dpy-30 does encode a protein. Additional evidence supporting this conclusion comes from the DNA sequence analysis of the two dpy-30 alleles, which had been classified as complete or nearly complete loss-of-function mutations by genetic analysis (Hsu and Meyer, 1994). The dpy-30(y130) mutation is a G to A transition at nucleotide 1015, resulting in a Gly to Arg missense change at amino acid position 85 in the predicted protein (Fig. 3). The dpy-30(y228) mutation is a C to T transition at nucleotide 925 that creates an amber stop codon at amino acid position 55 (Fig. 3). This finding is consistent with the genetic analysis demonstrating that y228 is partially suppressed by an amber tRNA suppressor. Molecular confirmation that y228 is an amber allele that disrupts the coding capacity of the dpy-30 gene provides strong evidence that y228 is a null allele of dpy-30.

DPY-30 is a ubiquitous nuclear protein whose expression pattern is unaffected by mutations that disrupt dosage compensation

Activation of the hermaphrodite mode of dosage compensation requires the sex-specific localization of the DPY-27 protein to the X chromosomes in XX animals (Chuang et al., 1994). Immunofluorescence experiments have demonstrated that the DPY-30 protein is one of the components essential for the proper localization of DPY-27 to X (P.-T. Chuang and B. J. Meyer, unpublished). To gain insight into the role of DPY-30 in this activation step and to explore the temporal and spatial pattern of DPY-30 expression in wild-type XX and XO animals as well as XX animals harboring mutations that disrupt dosage compensation, we performed immunolocalization studies using polyclonal antibodies raised against DPY-30 and confocal microscopy to detect the in situ antibody staining pattern (Materials and methods). The distribution of DPY-30
was visualized by indirect immunofluorescence with fluorescein-labeled antibodies, and the morphology of interphase nuclei was visualized by staining with the nucleic acid-intercalating molecule propidium iodide (PI). Comparison of the DPY-30 staining pattern (Fig. 4A) with that of the PI pattern (Fig. 4B) reveals that the two patterns are identical in embryos of all ages (Fig. 4C), demonstrating that DPY-30 is localized to the nucleus, consistent with its role in the regulation of X-linked gene expression. In contrast to DPY-27, DPY-30 is not associated with the X chromosome, or with any of the autosomes (D. Albertson, personal communication). This diffuse nuclear staining pattern was also observed in embryos (Fig. 5A), hermaphrodite oocytes, larvae and adults (data not shown) using epifluorescence microscopy, a finding consistent with the ubiquitous expression of dpy-30 mRNA and a possible requirement for dpy-30 throughout development.

Two lines of evidence indicate that the observed staining pattern accurately reflects the normal distribution of DPY-30 in the embryo. First, preimmune serum does not stain wild-type embryos (data not shown). Second, DPY-30 staining is eliminated in embryos homozygous for the null mutation dpy-30(y228) (Fig. 5C,D). To rule out the possibility that the reduced staining in y228 embryos is due to a general reduction in protein synthesis in these dying embryos, we also stained them with another antibody specific to the nuclear protein UNC-86 and obtained staining patterns similar to those of wild-type embryos (data not shown). These results indicate that the disappearance of DPY-30 antibody staining in homozygous y228 embryos reflects a true reduction in the level of DPY-30 epitopes, possibly due to the instability of either the dpy-30 mRNA or the truncated protein.

Since dosage compensation is a sex-specific process essential for the viability of only XX animals, we asked whether the pattern of DPY-30 expression differs in XX and XO animals, as does that of DPY-27. Previous genetic studies had demonstrated a role for dpy-30 in XO animals and previous molecular studies indicated that the dpy-30 transcript is expressed throughout development in both sexes, suggesting that DPY-30 expression is unlikely to be regulated in a sex-specific fashion. To identify XO embryos from a mixed population of XX and XO embryos, we took advantage of the sex-specific expression pattern of a xol-1::lacZ transgene, in which lacZ is under the control of the xol-1 promoter. XO transgenic embryos carrying an integrated version of this transgene express lacZ at a much higher level than their XX siblings (Rhind et al., 1995). XX and XO embryos from a strain homozygous for both the xol-1::lacZ transgene and a him-8 mutation, which increases X chromosome nondisjunction and thus the number of XO animals, were double stained with an anti-β-galactosidase monoclonal antibody and the DPY-30 antibody. The DPY-30 staining pattern in embryos identified as XO by virtue of their intense staining with anti-β-galactosidase antibody is identical to that of equivalently staged XX embryos (Fig. 6A-C). Thus, at the level of protein expression and localization, DPY-30 does not appear to be sex-specifically regulated.

The lack of sex-specific regulation of DPY-30 suggested the possibility that DPY-30 is a constitutively expressed protein whose pattern of expression is not influenced by other genes involved in dosage compensation. To test this hypothesis, we examined the DPY-30 staining pattern in mutant embryos defective in one of the several genes involved in the dosage sequence. The symbol above nucleotide 925 and the * symbol above nucleotide y130 mutation is a G to A transition that changes the Gly codon (GGA) to an Arg codon (AGA). Insertion of either the dpy-30 mRNA or the truncated protein.

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Fig. 3. DNA sequence of dpy-30. Shown is the 1363 nucleotide genomic region of pDH38 that encodes dpy-30. The full-length dpy-30 transcript, deduced from cDNA clones, is represented by DNA sequence listed in capital letters. The amino acid sequence of the predicted dpy-30 transcript is expressed throughout development in both sexes, suggesting that DPY-30 expression is unlikely to be regulated in a sex-specific fashion. To identify XO embryos from a mixed population of XX and XO embryos, we took advantage of the sex-specific expression pattern of a xol-1::lacZ transgene, in which lacZ is under the control of the xol-1 promoter. XO transgenic embryos carrying an integrated version of this transgene express lacZ at a much higher level than their XX siblings (Rhind et al., 1995). XX and XO embryos from a strain homozygous for both the xol-1::lacZ transgene and a him-8 mutation, which increases X chromosome nondisjunction and thus the number of XO animals, were double stained with an anti-β-galactosidase monoclonal antibody and the DPY-30 antibody. The DPY-30 staining pattern in embryos identified as XO by virtue of their intense staining with anti-β-galactosidase antibody is identical to that of equivalently staged XX embryos (Fig. 6A-C). Thus, at the level of protein expression and localization, DPY-30 does not appear to be sex-specifically regulated.

The lack of sex-specific regulation of DPY-30 suggested the possibility that DPY-30 is a constitutively expressed protein whose pattern of expression is not influenced by other genes involved in dosage compensation. To test this hypothesis, we examined the DPY-30 staining pattern in mutant embryos defective in one of the several genes involved in the dosage

Fig. 3. DNA sequence of dpy-30. Shown is the 1363 nucleotide genomic region of pDH38 that encodes dpy-30. The full-length dpy-30 transcript, deduced from cDNA clones, is represented by DNA sequence listed in capital letters. The amino acid sequence of the predicted dpy-30 translation product is listed below the DNA sequence. The symbol above nucleotide 925 and the * symbol above nucleotide y130 mark the positions of the nucleotide changes associated with dpy-30(y228) and dpy-30(y130), respectively. The y228 lesion is a C to T transition that changes the glutamine codon (CAG) to an amber codon (TAG). The y130 mutation is a G to A transition that changes the Gly codon (GGA) to an Arg codon (AGA). Insertion mutations at the designated BclI (nucleotide 689) and EcoRI (nucleotide 1057) restriction sites alter the wild-type dpy-30 coding region after codons 3 and 99, respectively, and eliminate rescuing activity. The GenBank accession number for dpy-30 is U21302.
compensation process. We found that XX embryos carrying null mutations in sdc-1, sdc-2, sdc-3, dpy-21, dpy-26, dpy-27
or dpy-28, have a DPY-30 staining pattern that is indistin-
guishable from the pattern observed in wild-type embryos,
indicating that DPY-30 expression does not depend on any of
the known dosage compensation genes or master regulatory
genes involved in the coordinate control of sex determination
and dosage compensation (Fig. 5E-H and data not shown).

Transcription of dpy-30 in the maternal germ line or
during gastrulation is sufficient to rescue both dpy-
30 XX and XO-specific phenotypes

To assess when transcription of dpy-30 is essential to guarantee
proportion X chromosome expression, we constructed a heat-
inducible dpy-30 transgene that allowed us to supply wild-type
dpy-30 transcripts to mutant animals at different times in develop-
ment. The heat-inducible transgene was created by fusing
the promoter of the C. elegans small heat-shock protein hsp16-
41 (A. Fire, personal communication) to a wild-type copy of
the dpy-30 coding region. Animals carrying an integrated copy
of the hsp16-41::dpy-30 fusion (ys15) were then used to determine the time during which dpy-30 transcription could rescue the XX-specific lethality and the XO-specific morphological defects caused by dpy-30 mutations. These experiments allowed us to determine that dpy-30 transcription in the mother
or in the embryo prior to the end of gastrulation is sufficient
to fully rescue both the XX- and XO-specific defects caused by dpy-30 mutations.

We determined the temporal requirements for dpy-30 tran-
scription to rescue the XX-specific lethality by applying heat
shocks to synchronized populations of dpy-30(ys228) XX
embryos at different times and scoring hermaphrodite viability
(Materials and methods). Fig. 7 shows that there are two distinct
periods when dpy-30 transcription can rescue the XX-specific lethality. Half hour heat shocks applied between 3 and 10 hours prior to first cleavage, or between 0.5 and 4 hours after first
 cleavage rescue greater than 80% of dpy-30 XX animals. In
contrast, heat shocks applied at 15.5 hours prior to first cleavage
or between 6 hours of development and hatching fail to produce
substantial numbers of viable hermaphrodites. Curiously, Fig.
7 also shows that heat shocks centered around the time of fer-
tilization (~0.5 hours) fail to rescue. Since heat shocks imme-
diately prior to or subsequent to fertilization prevent lethality,
it seems unlikely that the failure of heat shocks during fertil-
ization to overcome lethality reflects a time when dpy-30 tran-
scription cannot rescue XX dpy-30 animals. Moreover, the heat-
shock treatment itself cannot be responsible for the lack of
viability (Rhind et al., 1995). The most plausible explanation is
that this failure to rescue reflects a period in early embryonic
development when the heat-shock promoter is not sufficiently
active to generate the required level of dpy-30 transcripts.

The heat-shock experiments with dpy-30(ys15; dpy-30(ys228) XX
animals reveal a strong correlation between rescue of the XX-
specific lethality and rescue of the XX-specific morphological
defects (Dpy, Egl and Pvu) caused by the dosage compensa-
tion defect (Fig. 7). Cohorts with the highest viability also
contained the highest proportion of wild-type hermaphrodites.
In contrast, when rescue of viability was low, XX survivors
appeared variably Dpy, Egl and Pvu. This close parallel
between the rescue of viability and morphology is consistent
with studies of other C. elegans dosage compensation mutants,
suggesting that the penetrance and severity of the Dpy, Egl and
Pvu defects observed reflects the severity of their dosage com-
ensation defect (Plenefisch et al., 1989). These heat-shock
studies show that single heat shocks applied during gastrula-
tion result in the production of completely wild-type her-
maphrodites. We find no evidence for additional temporal
requirements for heat-shock-induced dpy-30 transcription.

We also determined the temporal requirement for dpy-30
transcription to rescue the dpy-30 XO-specific morphological
defects by heat shocking yIs15; dpy-30(y228) him-5 animals.
The him-5(e1490) mutation causes X chromosome nondis-
junction, resulting in the production of approximately 30% XO
male progeny. Males produced in these experiments were
scored for rescue of uncoordinated movement, small body size
and malformed tail structures associated with dpy-30 XO
males. Fig. 7 shows that heat shocks applied to the mother
during germline development or to the embryo during 2 to 6
hours of development is sufficient to rescue the dpy-30 XO-
specific morphological defects. As the case for rescue of
XX-specific lethality, little or no rescue of the XO defects is
apparent in the hours centered around fertilization.

dpy-30 is probably cotranscribed with rnp-1, a gene
encoding an unusual RNA-binding protein

The C. elegans genome contains many gene clusters that are
cotranscribed, as in bacterial operons. The individual message
for each gene is subsequently formed by polyadenylation and
trans-splicing (Blumenthal, 1995). Genes that comprise these
poly-cistronic transcription units share two major characteristics:
they are in close proximity, typically separated by a distance
of approximately 100 bp, and they have an unusual trans-splicing
pattern. The upstream gene is generally trans-spliced to the 22
nucleotide SL1 trans-spliced leader sequence. The downstream
gene is trans-spliced to the similar, but distinct, SL2 leader
sequence. While the functional significance of these poly-
cistronic transcription units is unknown, their existence suggests
the possibility that the genes in these C. elegans operons could
be functionally related, as they are in prokaryotes.

While cloning the dpy-30 locus, we identified a new gene,
rnp-1, whose proximity to dpy-30 and trans-splicing pattern
suggested that it might be cotranscribed with dpy-30. Com-
parison of the DNA sequence from genomic DNA and a rnp-
1 cDNA revealed that the rnp-1 transcript starts within 100 bp
downstream of the dpy-30 polyadenylation site (Fig. 8A).
Fur-
thermore, an SL2 trans-spliced leader sequence is present at
the 5′ end of rnp-1, as determined by the isolation of rnp-1
cDNA clones using RT-PCR with an oligonucleotide comple-
mentary to SL2 and an oligonucleotide internal to rnp-1 (data
not shown). No product was obtained when an oligonucleotide
complementary to SL1 was used in similar experiments.
These observations, together with the fact that dpy-30 is an SL1-
accepting gene fulfill the criteria for genes known to be tran-
scribed in C. elegans operons, suggesting that dpy-30 and rnp-
1 may also be cotranscribed.

DNA sequence analysis revealed that the mature rnp-1 tran-
script encodes a 305 amino acid protein with two distinct
nucleic acid-binding domains. The first domain, encompassing
approximately the first 80 amino acids of RNP-1, has signifi-
cant similarity to the consensus RNA recognition motif (RRM)
declared by a large and diverse family of proteins that bind pre-
mRNA, mRNA, snRNA and pre-rRNA molecules (for review,
**Fig. 4.** Nuclear localization of DPY-30 in XX embryos. (A-C) Confocal images of a wild-type XX embryo stained with (A) anti-DPY-30 antibody (false green color) and (B) counterstained with the nucleic acid-intercalating dye propidium iodide (PI) (false red color). The merged image (C) (false yellow color), which was created by superimposing the images of A and B, shows that DPY-30 is diffusely distributed throughout the entire nucleus of each cell in this embryo. Scale bar, 10 μm.

**Fig. 5.** Expression of DPY-30 in wild-type and dosage compensation defective embryos. Fluorescent photomicrographs of XX embryos stained with anti-DPY-30 antibody (A,C,E,G) and costained with DAPI to visualize nuclei (B,D,F,H). The wild-type nuclear staining pattern of DPY-30 is not perturbed by mutations that disrupt dosage compensation. Scale bar, 10 μm.
see Bandziulis et al., 1989) (Fig. 8B,C). This motif, found in single or multiple copies, is thought to be essential for the RNA-binding activities of these proteins. The alignment of the RNP-1 sequence with the RRM consensus shows that the RRM of RNP-1 has most of the conserved amino acids within this domain. Of particular importance is the fact that RNP-1 is nearly identical to the RRM of the cellular nucleic acid-binding protein (CNBP), seven copies of a similar motif are present in a seven stranded DNA (Covey, 1986). In addition, one or two copies of this motif, and define a group of low molecular weight retroviral proteins capable of binding RNA or single-stranded DNA (Covey, 1986). In addition, seven copies of a similar motif are present in a cellular nucleic acid-binding protein (CNBP), which binds a sterol regulatory element (SRE) in a sequence and single strand-specific manner (Rajavashisth et al., 1989).

The presence of both the RRM and NBP-type zinc finger motif provide a strong indication that rnp-1 acts as an RNA-binding molecule. Among RNA-binding molecules, RNP-1 is unusual in juxtaposing these two distinct RNA-binding motifs in a single protein. A BLAST search of both the PIR and GenBank databases revealed no other proteins with both RNA-binding domains. Thus, rnp-1 appears to define a novel class of RNA-binding proteins.

**DISCUSSION**

**Multiple roles of DPY-30 in development**

*C. elegans* has adapted an evolutionarily conserved chromosome condensation mechanism to achieve dosage compensation
(Chuang et al., 1994). It uses the X-chromosome-specific DPY-27 chromosome condensation protein to reduce expression of the hermaphrodite X chromosomes (Chuang et al., 1994). dpv-30 plays a critical role in this process, since it is essential for the sex-specific localization of DPY-27 to the hermaphrodite X chromosomes (P.-T. Chuang and B. J. Meyer, unpublished). However, dpv-30 is unique among dosage compensation genes in that it plays a more general role in C. elegans development than solely regulating X chromosome expression in hermaphrodites. Previous phenotypic analysis of dpv-30 mutants exposed two major differences between dpv-30 and the other maternal-effect dosage compensation genes, dpv-26, dpv-27, and dpv-28. First, null mutations in the other genes kill most, but never all XX animals, suggesting that the absence of dosage compensation is not completely lethal (Plenefisch, et al. 1989). In contrast, null mutations in dpv-30 cause complete X-specific lethality, suggesting that other developmental processes are aberrant in animals lacking dpv-30 activity (Hsu and Meyer, 1994). Second, while the other dosage compensation genes are dispensable in XO animals, dpv-30 is required for the wild-type development of XO males. dpv-30 XO mutants have wild-type X chromosome expression, but have numerous abnormalities including uncoordinated movement, reduced mating efficiency as well as abnormal tail and body morphology (Hsu and Meyer 1994). These pleiotropic phenotypes highlight the other developmental processes that require dpv-30 activity. The potential involvement of DPY-30 in the morphology and behavior of XX animals is obscured by the severity of the phenotypes caused by the dosage compensation defects.

The molecular results described in this paper further distinguish dpv-30 from other dosage compensation genes and are consistent with an involvement of DPY-30 in multiple processes. dpv-30 encodes a novel protein that is ubiquitously expressed in the somatic and germline nuclei of both XX and XO animals throughout development. DPY-30 is not associated with the X chromosomes nor with any autosomes, indicating that DPY-30's role in dosage compensation is fundamentally different from that of DPY-27. Unlike DPY-27, expression of DPY-30 is not affected by mutations in the other dosage compensation genes or in the master regulatory genes that control both sex determination and dosage compensation. However, we have not yet ruled out the possibility that DPY-30 is subject to sex-specific post-translational modification that is important for its role in dosage compensation. The lack of sex specificity in expression and the lack of obvious regulation by other dosage compensation genes suggests that DPY-30 participates in the localization of DPY-27 to the X chromosome by affecting the activity of another gene more directly involved in X chromosome binding. Likely candidates for such genes are sdc-2 and sdc-3, which are also required for proper localization of DPY-27 to the X chromosome (P.-T. Chuang and B. J. Meyer, unpublished). We have further shown that rescue of both the XX-specific lethality and the XO-specific morphological defects have similar, if not identical, temporal requirements for dpv-30 transcription, suggesting that dpv-30 mediates its diverse roles in XX and XO animals by acting at a similar time in both sexes. DPY-30 might affect multiple functions in males and hermaphrodites through its involvement in a general process such as transcription, or instead through its effects on a small number of genes having multiple roles in development. dpv-30 is not only unusual among the C. elegans dosage compensation genes, it is also different from the Drosophila dosage compensation genes (msls) that are required to elevate expression of the male X chromosome. All msl genes bind the hyperactivated male X and appear to play a direct role in elevating transcription (Baker et al., 1994).

![Proposed dpv-30 operon](image_url)

**Fig. 8.** Proposed dpv-30 operon. (A) Genomic structure and splicing pattern of dpv-30 and rnp-1. Analysis of both genomic and cDNA clones for dpv-30 and rnp-1 revealed that the 5' end of the rnp-1 transcript lies within 100 base pairs of the 3' end of the dpv-30 transcript. dpv-30 is trans-spliced to the SL1 leader and rnp-1 is trans-spliced to the SL2 leader. Rectangles represent exons in the mature transcript, lines represent introns and shaded regions indicate coding regions within each transcript. The extent of each gene is indicated underneath the genomic structure. (B) Predicted 305 amino acid translation product of the rnp-1 gene, with the RNA recognition motif (RRM) underlined and the putative zinc finger domain indicated by bold lettering. The GenBank accession number for rnp-1 is U21302. (C) Amino acid sequence alignment of the rnp-1 RRM and the CNBP consensus (Bandzülis et al., 1989), with the shaded boxes identifying conserved regions. rnp-1 shares the greatest similarity to the consensus in the two most highly conserved subdomains of the RRM, RNP 1 and RNP 2, which are underlined. (D) Amino acid sequence alignment of the rnp-1 zinc finger with consensus sequences for the CNBP and NBP families of zinc-fingers. Shaded rectangles indicate the conserved cysteine and histidine residues in these families. Numbers to the right of the rnp-1 sequence indicate the amino acid positions in the protein.
Timing of dosage compensation

Temperature-shift experiments revealed that dpy-30 transcription is only required early, either maternally prior to fertilization or during the first few hours of embryogenesis, to completely rescue dpy-30 mutants. These findings indicate that dpy-30 has an essential role in initiation of the XX mode of dosage compensation. Other lines of evidence are consistent with a role for dpy-30 in the activation of dosage compensation and further demonstrate that dosage compensation must be functional early in embryogenesis to ensure the viability of XX animals. The DPY-30-dependent localization of DYP-27 to the hermaphrodite X chromosomes, a key step in the activation of dosage compensation, occurs around the 30-cell stage (Chuang et al., 1994), the time when dpy-30 transcription is essential. In addition, temperature-shift experiments using a heat-sensitive allele of the dpy-28 dosage compensation gene established a requirement for dpy-28 around mid-embryogenesis (following gastrulation) for the viability of XX animals (Plenefisch et al., 1989). Finally, transcription of xol-1, a master switch gene whose activity state determines whether an embryo develops into a male or hermaphrodite, is required during gastrulation (2-4 hours following fertilization) to set the XO mode of dosage compensation (Rhind et al., 1995). Transcription of xol-1 in XX animals at the same early time causes nearly complete lethality because the hermaphrodite mode of dosage compensation cannot be activated. These results indicate that dosage compensation is functional soon after activation of zygotic transcription, which is thought to occur between the 8- and 16-cell stages (Edgar et al., 1994).

A question that remains unanswered by current studies is whether dosage compensation is required at any point after embryogenesis for wild-type development. Our studies indicate that dpy-30 mRNA and protein are present throughout the lifetime of XX animals and raise the issue of whether dpy-30 plays a role in dosage compensation throughout development. Biochemical studies have demonstrated that X-linked transcript levels are equalized between adult XO and XX animals, indicating that dosage compensation is active later in development (Meyer and Casson, 1986). Moreover, the sex-specific localization of DPY-27 to X persists throughout the lifetime of hermaphrodites (Chuang et al., 1994). This localization may require the continuous participation of DPY-30, since XX larvae and adults that lack wild-type DPY-30 fail to localize DPY-27 to X (P.-T. Chuang and B. J. Meyer, unpublished). Our results indicate that embryogenesis is a critical stage during which dosage compensation is essential for viability and raise the possibility that while dosage compensation occurs during later stages of development, it may not be essential for health and viability.

A dpy-30 operon

The dpy-30 and rnp-1 genes have all the properties common to C. elegans genes that are cotranscribed in an operon (for review see Blumenthal, 1995). The rnp-1 locus encodes a putative RNA-binding protein that is unusual in having both an RNA recognition motif (RRM) common to many RNA-binding proteins, as well as a single zinc finger motif similar to those from the nucleic acid-binding proteins (NBPs) of retroviruses. While the relationship of rnp-1 to the dosage compensation process is unknown, the fact that it appears to be encoded in an operon with dpy-30 presents the possibility that the two genes are functionally related and that dosage compensation may require aspects of RNA metabolism. Alternatively, evolutionary selection pressure on C. elegans might have caused a compaction of the genome, resulting in the clustering and co-regulation of genes that are related simply by their need for similar levels of expression in similar tissues (see Blumenthal, 1995).

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