Dosage compensation is a regulatory mechanism to ensure that the level of expression of genes on the single X chromosome of *Drosophila* males equals the level attained from the two X chromosomes in females. This equalization, achieved by a twofold increase in the rate of X-linked gene transcription in males relative to females, has been observed for a wide variety of genes with promoters of different strengths, in many cell types, and at different developmental stages. For this reason, the study of dosage compensation may provide valuable insights into the mechanisms that regulate levels of transcription.

Five genes involved in dosage compensation have been identified based on the male-specific lethality of their loss-of-function alleles (26). The products of these genes, collectively referred to as MSL proteins, co-localize to the male X chromosome, a chromosome that is also highly enriched with histone H4 acetylated at lysine 16 (6, 41). Since all eukaryotes acetylate the histones, it has been correlated directly with the establishment and regulation of transcription (reviewed in reference 28). It is likely that the MSL complex mediates its effect, at least in part, through histone acetylation. Indeed, the most recent MSL to be discovered is MOF (for “male absent on the first”), a protein with homology to acetyltransferases of the MYST family (8, 18, 33). Unlike the other members of the MSL complex, MOF can be specifically associated with the X chromosome in males (25).

Another protein component of the complex is MLE (for “RNA on the X 1 and 2”) that encodes RNAs with no apparent open reading frames (1, 27). This speculation has been reinforced, if not validated, by the recent discovery of two other forms of RNA (34). This RNA is found only in males, and its presence suggests that the complex may interact with either nascent or mature RNA (34). The MSL complex is responsible for the specific chromatin modification characteristic of the X chromosome in *Drosophila* males.

In *Drosophila*, dosage compensation—the equalization of most X-linked gene products in males and females—is achieved by a twofold enhancement of the level of transcription of the X chromosome in males relative to each X chromosome in females. A complex consisting of at least five gene products preferentially binds the X chromosome at numerous sites in males and results in a significant increase in the presence of a specific histone isoform, histone 4 acetylated at lysine 16. Recently, RNA transcripts (roX1 and roX2) encoded by two different genes have also been found associated with the X chromosome in males. We have partially purified a complex containing MSL1, -2, and -3, MOF, MLE, and roX2 RNA and demonstrated that it exclusively acetylates H4 at lysine 16 on nucleosomal substrates. These results demonstrate that the MSL complex is responsible for the specific chromatin modification characteristic of the X chromosome in *Drosophila* males.

In this paper, we report the initial functional characterization of the MSL complex. We demonstrate that all of the MSL proteins are associated in a complex that also contains a roX RNA. We also show that the complex requires MOF to acetylate H4 specifically on lysine 16, the isoform of H4 that colocalizes with the MSLs on the male X chromosome.

**MATERIALS AND METHODS**

**Antisera.** 12CA5 anti-HA monoclonal antibody was purchased from Boehringer Mannheim, and M2 Flag reagents were purchased from Sigma. Rabbit anti-H4Ac16 antibodies were generously provided by Brian Turner. Anti-MSL antibodies were raised against various fragments of MSL proteins fused to GST as follows: rabbit anti-MSL1 (amino acids [aa] 423 to 1029), guinea pig anti-MSL2 (aa 78 to 529), rabbit anti-MSL3 (full length), rabbit anti-MOF (aa 748 to 827), and rabbit anti-MLE (aa 1 to 359). Secondary antisera were purchased from Jackson Immunoresearch. Indirect-immunofluorescence images were collected with a Bio-Rad confocal microscope and false colored. Horseradish peroxidase-conjugated secondary antibodies were used for Western analysis and detected with enhanced chemiluminescence reagents (Amersham).

**Transfection and cell culture.** Schneider 2 (S2) cells were grown in SFX serum-free medium (HyClone). The calcium phosphate method of transfection was carried out as previously described (14). cDNAs were cloned into modified versions of the pMy/Hy vector (23) to allow for tagging with hemagglutinin (HA) or Flag epitopes. Stable transformants were selected by hygromycin in medium supplemented with 10% fetal calf serum, msl2-HA, mof-HA, and mof-HA encode full-length proteins with an extra 2 aa at the N terminus (Met-Ser-Glu) in place of the start Met and the HA epitope at the C terminus. msl3-Flag encodes full-length MSL3 with no additional amino acids at the amino terminus and the Flag epitope at the C terminus. Further details of the cloning or construction of the HA and Flag vectors will be provided on request.

**Preparation of nuclear extracts.** Cells were grown to a density of 3 × 10^6 cells/ml in 500-ml spinner flasks (Wheaton), pelleted at 750 × g for 5 min, washed in 200 ml of cell wash buffer (10 mM HEPES [pH 7.4], 140 mM NaCl), and resuspended in 80 ml of lysis buffer (20 mM HEPES [pH 7.4], 3 mM MgCl₂, 0.1% Triton X-100, 1 mM dithiothreitol [DTT], 0.5 mM phenylmethylsulfonyl fluoride [PMSF]). Cells were homogenized with 30 to 40 strokes of a Dounce homogenizer (Wheaton), post-nuclear clearance, 0.0035 to 0.0055 in., and the nuclei were pelleted at 2,000 × g for 5 min. The nuclei were washed once in lysis buffer and used for one of two extraction protocols, salt extraction or sonication. For salt extraction, nuclei were resuspended in 2 ml of extraction buffer (20 mM HEPES [pH 7.4], 10% glycerol, 0.35 M NaCl, 1 mM MgCl₂, 0.1% Triton X-100, 1 mM DTT, 0.5 mM PMSF) and rocked at 4°C for 1 h. Following centrifugation at 15,000 × g for 5 min, the supernatant was used for immunoprecipitation. For
 were washed 6 times in immunoprecipitation wash buffer (20 mM HEPES, 50 mM NaCl, 0.1% Triton X-100, 1 mM DTT, 0.5 mM PMFS, 5 U of RNasin [Promega] per ml) and sonicated on ice for three rounds of 10 s (output 4, 50% duty cycle [Branson Sonifier]) with 1-min rests between rounds. Nuclei were then pelleted for 5 min at 15,000 x g, and the supernatant was used for immunoprecipitation.

Immunoprecipitation. Crude antiserum (2 μl) or monoclonal antibody (20 μg) was bound to 10 μl of protein A agarose (BRL Gibco) for 1 h. Antibody-labeled beads were washed and incubated with 250 μl of salt extracts for 3 h. The beads were washed 6 times in immunoprecipitation wash buffer (20 mM HEPES [pH 7.4], 10% glycerol, 0.1 M NaCl, 1 mM MgCl₂, 0.1% Triton X-100, 1 mM DTT, 0.5 mM PMFS).

Two-step immunoprecipitation. M2 Flag agarose (40 μl) was added to 1 ml of nuclear sonicate (from MSL3-Flag expressing cells) and rocked at 4°C for 1 h. The beads were then washed five times with 1 ml of sonication buffer without RNasin. Bound proteins were eluted by the addition of 200 μl of 0.2 mg/ml Flag peptide (Asp-Tyr-Lys-Asp-Asp-Asp-Asp-Lys) and rocking for 20 min at 4°C. The eluate was passed through microspin filters (Pierce), and the flowthrough was applied to 10 μl of PI- or MSL1-bound agarose beads (see above) that had been preincubated for 30 min with 5 U of RNasin per ml in sonication buffer. After a 1-h incubation, the beads were washed six times with 400 μl of sonication buffer.

Histone acetyltransferase assays. (i) Purification of substrates. Drosophila histones (with H1) were extracted from S2 cell nuclei with 0.4 N H₂SO₄, pre-incubated for 30 min with 5 mM MgCl₂, and loaded on a 15 to 35% sucrose gradient (10 mM HEPES [pH 7.4], 10% glycerol, 0.1 M NaCl, 1 mM MgCl₂, 0.1% Triton X-100, 1 mM DTT, 0.5 mM PMFS).

(ii) Assays. A recombinant MOF fragment (aa 518 to 827) was expressed in Escherichia coli as a His-tagged fusion protein in pET11e (Novagen). Recombinant yeast proteins Gen5p and Esa1p were described previously (9, 37). A 5-μg portion of Drosophila histones was incubated with enzyme-0.2 μl of 3H-labeled acetyl coenzyme A (acytetyl-CoA) (7.2 Ci/mmol; Amersham)–1 mM DTT–1 mM PMFS–10% glycerol–50 mM Tris (pH 7.5) in a 50 μl-volume at room temperature (ca. 22°C) for 20 min. Aliquots were separated by SDS-PAGE, stained with Coomassie blue, and fluorographed with Amplify (Pharmacia) to identify which histones were labeled. Acetyltransferase assays with recombinant proteins were performed as above with the following changes: 2 μg of Drosophila nucleosome was incubated with 10 μl of protein A beads-immune complexes and 0.2 μCi of 3H AcCoA in a volume of 30 μl for 45 min with occasional mixing. The beads were spun down, and the supernatant was used for SDS-PAGE and fluorography.

Acid-urea gels. Aliquots of acetylation reaction mixtures were incubated with HCl (0.2 M final concentration) for 10 min on ice, trichloroacetic acid was added to 20% (wt/vol), and the mixture was incubated on ice overnight. After being washed with acetone, the pellets were resuspended in sample buffer and processed for discontinuous acid-urea gel electrophoresis as described previously (7). The gel was stained with Coomassie blue and processed for fluorography. Acid-extracted histones from S2 cells treated with butyrate were used as electrophoretic markers (13).

Determination of lysine specificity. The acid-urea gel was blotted to a polyvinylidene difluoride (PVDF) membrane (Millipore P-SQ) in 0.7 M acetic acid at 1 A for 2 h. The blot was stained briefly with Coomassie blue. The monoacetylated band was cut out and treated for deblocking and microsequencing as previously described (38).

RESULTS

MOF is a histone acetyltransferase. We expressed a cDNA fragment containing the putative MOF catalytic domain (aa 518 to 827) and determined that the recombinant peptide can acetylate Drosophila histones with a preference for histone H4 (Fig. 1). This pattern is similar to that for a related yeast protein, Esa1p (37), but different from that of the Gcn5-related enzymes (9). Since we were unsuccessful at expressing active full-length MOF, we proceeded to isolate MOF as a component of a partially purified MSL complex.

Partial purification of the MSL complex. We have chosen to use tissue culture cells for the initial characterization of the MSL complex. S2 cells are male, based on the following criteria: they do not express the Std (Sex- lethal) gene product, which is necessary for female differentiation (35), and they express MSL2, a limiting component of the dosage compensation machinery whose synthesis is prevented by SXL (4, 5, 21, 22). S2 cells can be stably transfected, allowing the use of commercially available antibodies recognizing epitope tags. Transient transfection of S2 cells with MSL2 tagged at its carboxy terminus with the HA epitope revealed that the localization of the HA epitope is coincident with the location of endogenous MOF (Fig. 2A). After selection with hygromycin, most cells exhibit HA staining on the male X chromosome, the location of which is revealed by antibodies to H4Ac16 (Fig. 2B).

Immunoprecipitation of nuclear extracts from MSL2-HA cells with the 12CA5 (anti-HA) antiserum resulted in the same proteins as those obtained from S2 cells with an MSL1 antiserum (Fig. 3). As can be seen from comparisons of silver-stained gels and Western blots, major bands seen by silver staining correspond to known MSLs, with the exception of MLE, which is difficult to detect under these conditions (see below).

In salivary gland nuclei, MLE is released from the male X chromosome with RNase treatment (34). Furthermore, the roX1 and roX2 RNAs are found along the X chromosome with a distribution that mimics that of the MSL complex (15). Therefore, we wished to determine if we could obtain a partially purified complex containing MLE and a roX RNA and whether the presence of either of these components depended on the other.

First, we developed “RNA-friendly” conditions to increase our chances of purifying MLE and roX RNA-containing complex. Our method involved a cell line expressing Flag-tagged MSL3 and sonication under low-salt conditions, immunopre-
cipitation with Flag antibodies followed by peptide elution, and a second immunoprecipitation with an MSL antibody or with the corresponding preimmune serum. By using this two-step procedure, we can detect a faint band by silver staining that corresponds to MLE protein (Fig. 4A). Clear enrichment of MLE was seen in the MSL1 immunoprecipitate relative to the preimmune serum (Fig. 4B). However, following a brief treatment with 0.4 M NaCl, the MLE levels were significantly reduced.

To determine if roX RNAs are expressed in S2 cells, we performed Northern blot analysis and observed that roX2, but not roX1, was expressed in these cells (Fig. 5A), consistent with the observation that roX1 is dispensable in flies (27). The size of the major roX2 transcript observed by Northern analysis was ca. 600 nucleotides. This size is consistent with the results reported by Amrein and Axel (1) in their Fig. 2, which demonstrate that roX2 transcripts migrate slightly faster than the 600-nucleotide rp49 mRNA (29). To test if roX2 RNA is

FIG. 2. Epitope-tagged MSL2 colocalizes with MOF and histone H4 acetylated at lysine 16 (H4Ac16) on the X chromosome in S2 cells. (A) Cells were transfected with a construct expressing MSL2 fused to the HA epitope. Tagged MSL2 is detected with the 12CA5 monoclonal antibody, while endogenous MOF is detected with a rabbit polyclonal antibody to the C terminus of MOF. The top panel shows that a small fraction of the cells contain transfected MSL2-HA, which is seen colocalizing with endogenous MOF. The bottom panel shows an enlargement of a nucleus where both antibodies can be seen painting a chromosome that traverses the nucleus. (B) Stable cell lines were selected with hygromycin, and a line that expressed a low level of basal (uninduced) expression was chosen for further study. In this line, basal levels of MSL2-HA can be detected along the length of the X chromosome, coincident with H4Ac16 and the other MSLs (not shown). Occasional areas of nonoverlap may reflect the presence of partial or nonfunctional complexes.
present in the MLE-containing immunoprecipitates, RNA was extracted from the immunoprecipitation pellets and RT-PCR was performed with roX2-specific primers in the linear range. The results show a clear enrichment of roX2 RNA in the immune over the preimmune serum precipitates (Fig. 5B).

The MSL complex specifically acetylates lysine 16 of histone H4. When MSL-containing immunoprecipitates were incubated with nucleosomal substrates, significant acetyltransferase activity toward histone H4 was detected (Fig. 6A). MSL1 immunoprecipitates from S2 nuclear extracts and 12CA5 immunoprecipitates from MSL2-HA nuclear extracts contain H4-specific acetyltransferase activity, while control immunoglobulin G or 12CA5 immunoprecipitates from S2 cells do not.

To demonstrate that the acetyltransferase activity of the MSL complex is ascribable to MOF, we purified complexes containing either wild-type MOF or a protein produced by the mutant allele mof1 (18). This allele is a point mutation resulting in a glycine-to-glutamic acid replacement at the most highly conserved residue of the acetyl-CoA binding domain (G691E). We overexpressed wild-type MOF-HA or G691E MOF-HA in S2 cells and immunoprecipitated them with anti-HA antibodies to obtain complexes with only transfected MOF fusion proteins. Immunoprecipitates from G691E cells have markedly reduced acetylation, consistent with the conclusion that MOF is the sole acetyltransferase in the MSL complex (Fig. 6B).

Given the specificity of the MSL complex toward H4, we wished to determine which particular lysines were acetylated. When acetylated histones were separated by acid-urea gel electrophoresis, predominantly monoacetylated H4 was created (Fig. 7, inset). A similar acid-urea gel was blotted to PVDF, and the mono-acetylated band was subjected to microsequenc-

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**FIG. 3.** Major immunoprecipitated proteins detected by silver staining correspond to known MSLs. The 12CA5 antibody (HA) generates the same set of proteins from MSL2-HA (M2HA) nuclear extracts that the MSL1 antiserum generates from S2 cell extracts; differences in stoichiometry are ascribable to the overexpression of HA-tagged MSL2. This set of proteins is absent from HA or control immunoglobulin G (PI) precipitates from (untransfected) S2 cells. Western analysis of the MSL1 immunoprecipitate shows that the major silver-stained bands correspond to known MSLs. Rabbit anti-MSL1 serum detects a 170-kDa band, guinea pig anti-MSL2 detects a 135-kDa band, and rabbit anti-MSL3 detects a 58-kDa doublet of bands. Molecular masses are indicated in kilodaltons (KDa).

**FIG. 4.** MLE association with the MSL complex. Significant levels of MLE are detected with other MSLs when immunoprecipitations are performed under low-salt conditions (see Materials and Methods). MSL complexes were eluted from M2-Flag agarose and subjected to a second immunoprecipitation with anti-MSL1. (A) A silver-stained protein is visible between MSL1 and MSL2/MOF. Two concentrations (1× and 5×) of a stringently washed anti-MLE immunoprecipitate (IP) were loaded on the same gel; correlation between silver staining and Western staining intensities as well as comigration by SDS-PAGE, confirms that this band is MLE. (B) Comparison of low-salt (LS) and high-salt (HS) washing conditions reveals a salt-sensitive association of MLE with the other MSLs. As seen by Western analysis, significant levels of MLE are released from immunoprecipitates when incubated with high-salt buffers but not low-salt buffers; MSL1 levels are unaffected. A preimmune control (PI) was washed under low-salt conditions and reveals a low level of contaminating MSLs in these preparations.
Counts were found at lysine 16, while other potential acetylation sites (at position 5, 8, or 12) were unlabeled (Fig. 7). This result provides a causative link between the presence of histone H4 acetylated at lysine 16 and the MSL complex on the X chromosome in *Drosophila* males.

**DISCUSSION**

The last few years have seen a remarkable increase in our knowledge of macromolecular complexes that modify chromatin and thereby modulate gene activity. These complexes interact with nucleosomal proteins, i.e., histones and in some cases with components of the preinitiation complex to regulate the level of transcription of a large number of different genes. At present, these multiprotein complexes can be placed into two broad categories: complexes that use the energy of ATP hydrolysis to alter nucleosomal conformation, and complexes that alter chromatin conformation via the modification of histone tails (reviewed in reference 45). A few examples of the former category are the RSC and SWI/SNF complexes in yeast (10, 11, 32), and the Brahma, NURF, CHRAC, and ACF complexes in *Drosophila* (19, 31, 40, 43). In these complexes the ATP requirement is ascribed to the activity of proteins which contain domains characteristic of helicases but which do not exhibit in vitro helicase activity. Complexes in the second category, including SAGA and NuA4 from yeast (17), are

A

**FIG. 5.** *roX2* is expressed in S2 cells and associates with the MSL proteins. (A) Northern analysis shows that both *roX* RNAs are expressed in adult *Drosophila* males (M) but not females (F); only *roX2* was detected in S2 cells. The filters were reprobed for *rp49* RNA as a loading control. (B) RNA was extracted from anti-MSL1 immunoprecipitates (M1) or the corresponding preimmune serum (PI) and subjected to RT-PCR. Agarose gel electrophoresis of PCR products and staining with ethidium bromide detected significant levels of *roX2* RNA in the anti-MSL1 immunoprecipitates, while a lower level of *roX2* RNA (1 to 2%) was detected in the preimmune serum. This level of contamination is consistent with the amount of contaminating MSL proteins observed in these immunoprecipitates (Fig. 4B). A kanamycin kinase transcript was used to monitor variation in processing of samples; comparable levels of this control RNA (Ctrl) were detected by RT-PCR in the preimmune and anti-MSL1 immunoprecipitates.

B

**FIG. 6.** The MSL complex acetylates nucleosomal H4 in a MOF-dependent manner. (A) Immunoprecipitates were assayed for acetyltransferase activity toward mononucleosomes and processed for SDS-PAGE and fluorography. Both MSL1 immunoprecipitates (IP) from S2 cells and 12CA5 immunoprecipitates (HA) from MSL2-HA-expressing cells acetylate nucleosomes specifically on histone H4. Preimmune serum (PI) or 12CA5 monoclonal antibody does not immunoprecipitate any histone acetyltransferase activity from S2 cells. (B) Acetyltransferase activity was also assayed from cells transfected with the HA-tagged *mof<sup>1</sup>* or HA-tagged *mof<sup>2</sup>* allele (Gly-to-Glu mutation at residue 691). Western analysis demonstrates that the extracts contain complexes with large amounts of HA-tagged MOF or MOF<sup>1</sup>. Histone acetyltransferase activity from MOF<sup>1</sup>-containing complexes is dramatically reduced.
thought to function by targeting specific histone acetyltransferases to their site of action for the activation of transcription (24, 42). As described in this paper, the MSL complex includes both an ATP-dependent helicase (MLE) and a histone acetyltransferase (MOF); it also contains one or more noncoding RNAs of unknown function (roX).

It is notable that S2 cells express roX2 but not roX1 RNA. This, however, is consistent with the observation that these two RNAs are functionally redundant (15, 27). The roX RNAs differ vastly in size (3.5 and 0.6 kb for roX1 and roX2, respectively) and lack extensive similarities, although a small region of unknown function was identified by Franke and Baker (15).

One function of roX RNAs may be to help maintain the association of MLE with the other MSLs. MLE is distantly related to the SWI2 and ISWI ATPases, but it has been shown to exhibit RNA/DNA helicase, ATPase, and single-stranded RNA-DNA binding activities in vitro (25). As previously mentioned, MLE can be released from larval salivary gland chromosomes by RNase treatment (34) and appears to associate rather weakly with the other protein components of the complex (13). Our ability to detect significant amounts of MLE in the complex is probably due to shortened processing time, low-ionic-strength buffers, and RNase-free conditions.

MOF is a member of the MYST family of histone acetyltransferases. Two other members of this family, Esa1p and Tip60, acetylate histones H2A, H3, and especially H4 when tested in vitro as fusion proteins or as catalytic fragments (12, 37, 46). The specificity of the Esa1p-containing NuA4 complex, with respect to the sites of histone H4 acetylation, is identical to that of the recombinant Esa1p: lysines 5, 8, and 12 and 16 (37). Recently, a protein (p80) has been identified in Tetrhydymena that acetylates nucleosomal H4 in an identical pattern (30). In contrast, the MSL complex shows a clear preference for lysine 16.

One proposed role of histone acetylation in transcription is to help promote transcription factor access to DNA for the purposes of initiating transcription. Gcn5p activity is responsible for the acetylation of histone H3 at the HIS3 promoter, which results in gene activation (24). Additionally, histone acetylation may facilitate elongation by RNA polymerase. This is supported by in vitro experiments with hyperacetylated histones and RNA polymerase III (39) and by the recent discovery that an RNA polymerase II-associated elongation complex contains acetyltransferase activity (44).

In Drosophila, the preponderance of histone H4 acetylated at lysine 16 is found associated with the MSL complex on the X chromosome in males (6, 41). Since the other isoforms of histone H4 are found in similar amounts on both male and female X chromosomes (41), one might infer that MSL complexes are targeted to chromatin already acetylated at other lysines; the complex then adds an acetate on lysine 16 to further increase transcription. Whether the increase in transcription occurs at initiation or elongation is currently under study.

Finally, it should be noted that the MSL complex of Drosophila enhances transcription not by orders of magnitude (as appears to be the case with some of the other chromatin remodeling complexes) but, on average, only by a factor of 2. Therefore, it may be a very good model to study how chromatin-remodeling complexes do, in fact, achieve particular levels of gene activity.

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