Cotranscriptional recruitment of the dosage compensation complex to X-linked target genes

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Dosage compensation is a process required to balance the expression of X-linked genes between males and females. In *Drosophila* this is achieved by targeting the dosage compensation complex or the male-specific lethal (MSL) complex to the male X chromosome. In order to study the mechanism of targeting, we have studied two X-chromosomal genes, *mof* and *CG3016*, using chromatin immunoprecipitation as well as immuno-FISH analysis on transgenic flies. We show that MSL complex recruitment requires the genes to be in a transcriptionally active state. MSL complex recruitment is reversible because blocking transcription severely reduces MSL binding to its target genes. Furthermore, targeting cues are found toward the 3' end of the gene and depend on the passage of the transcription machinery through the gene, whereby the type of promoter and the direction of transcription are dispensable. We propose a model of dynamic MSL complex binding to active genes based on exposed DNA target elements.

[**Keywords:** MSL; MOF; roX; transcription; dosage compensation; X chromosome]

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Sex determination often involves differentiation of a pair of chromosomes such that one sex retains the original two copies (homogametic), whereas in the other sex one of the homologous chromosomes degenerates almost completely with only the exception of a few sex-specific genes (heterogametic). The deleterious effects of aneuploidy creates the need for a compensatory process that equalizes the gene products between the two sexes (Charlesworth and Charlesworth 2005). In *Drosophila*, dosage compensation occurs by increasing transcription of most genes on the single male X chromosome to compensate for transcription of both X chromosomes in females. Genetic studies have identified five male-specific lethal (MSL) genes—*msl1*, *msl2*, *msl3*, *males absent on the first* (*mof*), and *maleless* (*mle*)—that are important for the regulation of dosage compensation. The products of these genes (collectively referred to as MSL genes), as well as two noncoding RNAs (roX1 and roX2), assemble in a large complex called the dosage compensation complex (DCC), or the MSL complex, which is targeted to hundreds of sites on the male X chromosome (Lucchesi et al. 2005; Mendjan and Akhtar 2006; Straub and Becker 2007).

One of the most intriguing features of the MSL com-

plex is its ability to recognize its targets on the male X chromosome. Studies that involved *P*-element-mediated insertions of X-chromosomal genes on autosomes and X-autosome translocations have shown that most X-linked genes were still targeted when inserted on an autosome. In contrast, autosomal genes translocated to the X remained untargeted (Fagegaltier and Baker 2004; Oh et al. 2004). Targeting therefore most likely happens on genes individually and may not require the presence of 35–40 so-called “entry sites” (Kelley et al. 1999).

Recent chromatin profiling studies have confirmed the preference of the MSL complex for individual genes. The MSLs were found to bind locally to genes with MSL occupancy predominantly at the 3' of genes (Alekseyenko et al. 2006; Gilfillan et al. 2006; Legube et al. 2006). Moreover, the affinity of MSL1 for its targets correlates with its dosage-compensated state (Gilfillan et al. 2006; Legube et al. 2006). Therefore, the affinity of the MSL complex might have evolved concurrent with the need for dosage compensation of its targets. This is in agreement with the findings that those genes strongly bound by MSL1 are more generally essential genes (Gilfillan et al. 2006).

Despite the large number of MSL targets recently identified by chromatin profiling studies, a universal targeting sequence motif has not been identified for the MSL complex. Also, the proposal that the MSL complex simply targets those genes on the X that are transcriptionally active does not seem to apply universally since there are many genes on the X that are transcriptionally active but

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not bound by the MSLs (Alekseyenko et al. 2006; Gilfillan et al. 2006; Legube et al. 2006). It seems conceivable that transcription is a prerequisite for target recognition but not sufficient by itself, because most target genes are transcribed, and those few genes that show differential expression are only bound when the gene is active (Alekseyenko et al. 2006). Although no universal targeting sequence has been found, several short degenerative sequences with some predictive power have been identified (Dahlsveen et al. 2006; Gilfillan et al. 2006). It is therefore possible that these degenerative sequences are only exposed when the gene is active and the chromatin is present in an open conformation.

An assumption of the above hypothesis would be the ability of the MSL complex to associate with genes only while transcription is ongoing, presumably at the stage of transcriptional elongation. Therefore, targeting cues may reside in the body of the gene as opposed to the original reports of upstream enhancer sequences present in dosage-compensated genes (for review, see Baker et al. 1994), but in agreement with recent MSL profiling studies (Smith et al. 2001; Alekseyenko et al. 2006; Gilfillan et al. 2006; Legube et al. 2006).

In order to gain better insights into the targeting mechanism, we studied two neighboring genes, *CG3016* and *mof*, for their targeting properties as they recently have shown to be targeted by the MSL complex in genome-wide analysis (Gilfillan et al. 2006; Legube et al. 2006). The *mof* gene encodes the histone acetyltransferase in the MSL complex that specifically acetylates Lys 16 of histone H4 (H4K16). Apart from the two *roX* RNA-encoding genes, *mof* is the only other known MSL member whose gene is located on the X chromosome. Interestingly, we find that transcriptional activation is required for targeting of the MSL complex and that polymerase passage through the gene is a prerequisite for target recognition. We also show that this is true for most genes on the X chromosome, as blocking transcription by α -amanitin treatment greatly reduced binding of the MSL complex to X-chromosomal genes. Finally, we show that in addition to transcription, targeting to the *mof* and *CG3016* genes most likely is a combination of degenerative DNA sequences toward the 3' end of these genes only to be recognized when the gene is transcriptionally active.

Results

MSL proteins are enriched on the region spanning the mof and CG3016 genes

In order to precisely map the binding pattern of MSLs on the *mof* and *CG3016* loci, we performed chromatin immunoprecipitation (ChIP) experiments in *Drosophila* Schneider cells (SL-2) using specific antibodies directed against MSL1, MSL3, and MOF. SL-2 cells are a male 16-h embryonic cell line with an MSL-binding pattern closely reflecting that of late-stage embryos (Alekseyenko et al. 2006). The immunoprecipitated material was analyzed by quantitative real-time PCR (Q-PCR) and

presented as the percentage recovery compared with input DNA (Fig. 1A). The *roX2* high-affinity site, which served as a positive control, showed highly enriched MSL binding (Fig. 1A, lane 15) as opposed to *gapdh* (located on chromosome 2R) (Fig. 1A, lane 14) and the *runt* gene (located on the X chromosome) (Fig. 1A, lane 13), but is dosage-compensated in an MSL-independent fashion (Gergen 1987; Smith et al. 2001). Binding was enriched on the body of the gene for both *CG3016* and *mof* for all three MSL proteins tested (Fig. 1, lanes 1–12). Unlike MSL1 and MSL3, there is also considerable binding of MOF to the promoter regions (see Fig. 1, lanes 1,5). Whether the binding of MOF to promoters reflects its general binding pattern on a genome-wide scale remains to be tested.

We next wished to test if a large genomic fragment including *mof* and most of *CG3016* was able to target the MSL complex ectopically when inserted on an autosomal location by means of *P*-element-mediated transformation. As is shown in Figure 1B, this 6.7-kb fragment (*mof*^{6.7}) showed robust MSL1 recruitment visualized by staining with an MSL1 antibody in combination with fluorescence in situ hybridization (immuno-FISH) to visualize the location of the transgene (see Materials and Methods). The presence of MSL1 and the direct visualization of targeting on polytene chromosomes enabled further detailed analysis of the nature of targeting to this locus.

Taken together, these results show that MSL1 protein is enriched on *mof* and *CG3016* genes and that this region is able to recruit the MSL complex autonomously on an autosomal location.

MSL1 is recruited to the mof gene in a transcription-dependent manner

In order to study targeting to individual genes, we first generated transgenic flies carrying the *mof* gene with its endogenous promoter sequences (*mof*^{3.5}). The *mof* gene showed ectopic MSL1 recruitment in almost all independent lines that were tested (Fig. 1C; Supplementary Table 1). Recently, the MSL-binding pattern to polytene chromosomes was suggested to reflect the distribution of active genes (Sass et al. 2003). Although the vast majority of genes bound by MSL1 are indeed active, several active genes on the X chromosome escape MSL1 binding (Alekseyenko et al. 2006; Gilfillan et al. 2006; Legube et al. 2006). Therefore, in order to address directly whether or not transcription is a prerequisite for targeting, we analyzed targeting of the MSL complex to the *mof* gene in the absence of the endogenous promoter sequences (*mof*^{2.4} Δ prom). Clearly, targeting to these lines was lost as we did not observe MSL1 binding on the transgene in any of the lines studied (Fig. 1D; Supplementary Table 1).

Although the results above suggest that the absence of transcription leads to the loss of MSL1 binding, it is possible that targeting sequences are present in the promoters of the respective genes as it was originally suggested for several genes (Baker et al. 1994). In order to address if the promoter requirement is due to a targeting signal

Figure 1. Transcriptional-dependent recruitment of MSL1 to the *mof* gene. (A) ChIP in SL-2 cells on the *CG3016* and *mof* genes with antibodies directed against MSL1, MSL3, and MOF. Levels of enrichment are determined by Q-PCR and depicted as the percentage recovery of input DNA. The position of the primer pairs is schematically shown by green arrows. (B–D) MSL-1 immunostaining (green) in combination with DNA-FISH (red) directed against the mini-white gene of a polytene chromosome carrying an autosomal 6.7-kb *P*-element insertion encompassing *mof* and the coding region of *CG3016* (B), or an autosomal *P*-element insertion of *mof* with its endogenous promoter (C) or without its endogenous promoter (D). Recruitment of MSL1 to the transgenes is indicated by an arrow. The position of the endogenous *mof* gene is shown by an asterisk. DNA was stained with Hoechst 3325 (blue).

embedded in the promoter sequence or solely due to transcriptional activation, *mof* was fused downstream from the *tubulin* promoter. The *tubulin* gene is located on chromosome 3, and therefore its promoter should not contain any X-chromosome-specific sequences. Interestingly, a *tubulin* promoter-driven *mof* transgene (*tub-mof*) also showed ectopic recruitment of MSL1 (Fig. 2A), as well as MSL3, MLE, and MOF (Supplementary Fig. 1). In contrast, *tubulin*-driven expression of the *pho* gene (Klymenko et al. 2006), encoded on the fourth chromosome, did not show any MSL1 recruitment, further confirming the specificity of MSL1 binding to the *mof* gene (Supplementary Fig. 2A; Supplementary Table 1).

In order to study transcription-dependent targeting at the same genomic position and thereby addressing any possible position effect variegation (PEV)-related effects, we also created a transcription on/off assay by using a previously developed system that makes use of FRT-flip recombinase-mediated excision of a *yellow* gene inserted between the *tubulin* promoter and the gene of interest, in our case the *mof* gene (Struhl and Basler 1993). When present, the *yellow* gene blocks transcription from the

tubulin promoter. This assay can therefore be used to study recruitment of MSL complex to the same location in the inactive and active states. Using this assay, we observed that *mof* was only recognized as a MSL1 target when transcription was allowed by excision of the *yellow* gene (Fig. 2B,C; Supplementary Table 1). We confirmed activation of transcription after excision of the *yellow* gene by rescuing the male-specific lethality caused by the absence of MOF (data not shown). Since *mof* driven by a tandem repeat of five GAL4-binding sites (*UAS-mof*) also showed targeting of MSL1 (Supplementary Fig. 2B; Supplementary Table 1), we conclude that targeting solely requires transcriptional activity irrespective of the promoter used. The determinant of targeting, therefore, most likely is to be found in the gene region and not in the promoter sequence. Furthermore, the observation that UAS-driven expression of *mof* without GAL4 activation also displayed MSL1 recruitment suggests that transcription is required but the levels of transcription are not important.

A limited number of sites on the X chromosome are defined as high-affinity sites because they have the abil-

Figure 2. The *mof* transgene driven by a *tubulin* promoter is a target for MSL1 and overcomes the requirement for MSL3. (A) Tubulin-driven *mof* gene. (B,C) Tubulin-driven *mof* gene (*tub > y+ > mof*) containing a *yellow* gene cassette between the promoter and the gene such that *mof* expression becomes inducible upon excision of the *yellow* gene cassette by expression of flippase. Recruitment of MSL1 before (B) and after (C) excision is shown. (D) Same as in A, however, in this case the transgenic line carrying *tub-mof* was crossed into the *msl-3⁰⁸³* mutant background by using females mutant for *msl-3⁰⁸³* expressing low levels of MSL2 from an *Hsp83* promoter. Recruitment of MSL1 to the *tub-mof* transgene is indicated by an arrow. The position of the endogenous *mof* gene is shown by an asterisk. Red cross (x) represents no transcription. DNA was stained with Hoechst 322 (blue).

ity to recruit partial complexes of MSL1 and MSL2 in the absence of the other MSLs (Kelley et al. 1999). Since *mof* is located in a cytological location of the previously mapped high-affinity site at position 5c (Lyman et al. 1997; Demakova et al. 2003), we tested by DNA-FISH if this site coincides with the location of *mof*. Since homozygous *msl-3⁰⁸³* male third instar larvae are rarely recovered, we used female flies expressing MSL2 in an *msl-3⁰⁸³* mutant background. These flies assemble partial complexes on the X chromosome and are routinely used to study high-affinity sites (Kelley et al. 1999; Kageyama et al. 2001; Dahlsveen et al. 2006). As shown in Figure 2D, the *mof* endogenous gene is found in very close proximity to the high-affinity site, but upon close examination appears to be not the site itself but just beside it (Fig. 2D, see asterisks). We verified this result by crossing the genomic construct containing the *mof* locus (*mof^{3.5}*) in an *msl-3⁰⁸³* background, which also showed no targeting (data not shown). Surprisingly, however, when the *tubulin*-driven *mof* transgene was crossed in an *msl-3⁰⁸³* background, we created what appeared to be a high-affinity site, as we could observe recruitment of MSL1 in the absence of MSL3 (Fig. 2D, arrow). We obtained similar results with this line in an *mle¹* mutant background, albeit with a substantially weaker MSL1 recruitment (data not shown).

Taken together, the results above show that the MSL complex is recruited to the *mof* gene in a transcription-dependent manner. Our results suggest that also on other regions of the X chromosomes, otherwise known as lower-affinity sites, partial MSL complexes are able to recognize their targets autonomously without MSL3 or MLE. We therefore propose that low-affinity sites may override the requirement for MSL3 or MLE if their expression is driven from a strong promoter such as *tubulin*. This is presumably due to the combinatorial effect of chromatin modifications associated with high levels of transcription that may cooperate to expose certain DNA target sequences to MSL1 and MSL2, which may not be exposed otherwise.

MSL complex targeting is compromised in α -amanitin-treated cells

Knowing that transcription is a prerequisite for binding of the MSL complex (Fig. 2B,C), we wished to test for the plasticity of the bound state. Somewhat conflicting hypotheses on the subject of timing and maintenance of targeting the MSL complex have been put forward recently. On one hand, the MSL complex is thought to be targeted in early development, presumably following the present transcriptional pattern, and maintained in the course of development irrespective of transcriptional changes (Alekseyenko et al. 2006; Legube et al. 2006). On the other hand, subtle developmental changes in the binding pattern of the MSL complex on polytene chromosomes and the presence of genes in early development (6-h embryos) that are bound by MSLs but do not correlate with transcriptional status, as judged by RNA Pol II chromatin-binding profiles (Gilfillan et al. 2006), suggest that differential binding takes place.

Since many MSL target genes are housekeeping genes that are continuously transcribed (Gilfillan et al. 2006; Legube et al. 2006), stable binding throughout development could be the rule with some differentially transcribed genes being the exception. In order to test this hypothesis, we blocked transcription by means of α -amanitin treatment in Schneider (SL-2) cells. We argued that if binding is maintained irrespective of transcriptional changes, genes should preserve their normal MSL-binding pattern when transcription is blocked. As expected, after a 20-h α -amanitin (Pol II inhibitor) treatment, transcript levels of most genes tested were significantly reduced compared with untreated control levels (Fig. 3A). Visually the cells showed normal morphology (data not shown), and the protein levels of MSL1 and MOF were comparable with the control situation (Fig. 3B). Furthermore, roX2 RNA levels in treated cells were mildly affected in comparison with the control situation, very likely reflecting a higher stability of this RNA (Fig. 3A). In addition to the *mof* gene, we tested four additional X-chromosomal genes (*UCP4A*, *SOCs16D*, *CG4061*, and *CG3016*) by ChIP. Interestingly, all the tested genes show a dramatic reduction of MSL binding following α -amanitin treatment (Fig. 3C). Intriguingly, MSL binding to the high-affinity sites such as *roX2* and *18D* was significantly less susceptible to α -amanitin treatment, presumably reflecting their ability to recruit the complex ectopically independently of transcription (Kelley et al. 1999; Kageyama et al. 2001; Oh et al. 2003, 2004). Consistent with these observations, we also found severely reduced staining of MSL1 on the X-chromosomal territory in α -amanitin-treated SL-2 cells in comparison with the control cells (Fig. 3D). These results suggest that MSL binding is reversible and depends directly on the transcriptional state of the targets. The rigid binding of MSLs therefore is not an embryonic irreversible preset condition, but most probably reflects the continuous transcriptional activity of its targets.

*Genic sequences contribute to MSL complex recruitment on the *mof* gene*

Although it is conceivable that targeting requires the transcription machinery to pass through the gene for a certain targeting signal to become exposed, it remains plausible that MSL binding is achieved by the recruitment of transcriptional activators to the promoter; e.g., in the scenario in which the MSL complex slides along the DNA pushed forward by transcribing polymerase (Gilfillan et al. 2006). For this purpose, we created a *mof* construct that has the *mof* endogenous promoter but in a reverse orientation (*mof^{3.5}*prom AS), such that transcription does not pass through the *mof* gene. As is shown in Figure 4A, this transgene is unable to recruit MSL1. Thus, transcription activation is not sufficient for targeting MSL1 to *mof*, instead transcription needs to run through the gene to reveal *mof* as an MSL target.

To date, it remains unknown what the targeting signal embodies. Several groups over the past years have attempted to identify a universal targeting sequence, but

Figure 3. Recruitment of MSL1 and MOF on X-chromosomal genes is severely affected upon blocking transcription by α -amanitin. (A) Q-RT-PCR analysis of expression levels normalized to mock (100%) after a 20-h α -amanitin treatment. (B) Western blot analysis from whole-cell extracts prepared from mock- or α -amanitin-treated cells. Western blots were probed with MSL1, MOF, and Tubulin antibodies as indicated. (C) ChIP in SL-2 cells with antibodies directed against MSL1, and MOF after 20-h mock or α -amanitin treatment. Levels of enrichment are determined by Q-PCR and depicted as the percentage recovery of input DNA. Enrichment on each gene was systematically tested using primer pairs located at the beginning, middle, and end of genes. Position of the PCR probes with respect to the transcription start site (+1) is indicated on the X-axis; for example, for UCP4A, the first primer pair is located 367 bp upstream of the transcription start site, and the second and third primer pairs are located 612 bp and 2860 bp downstream, respectively. (D) Immunofluorescence with MSL1 (red) and Lamin (green) in Schneider (SL-2) cells of mock- or α -amanitin-treated cells. DNA was stained with Hoechst 322 (blue).

despite these attempts, beside several short degenerative sequences with limited predictive power, no general binding sequence has been identified (Alekseyenko et al. 2006; Dahlsveen et al. 2006; Gilfillan et al. 2006; Legube

et al. 2006). It is possible that the predictive power of these sequences is limited because they need to be exposed in order to be recognized. It is conceivable, therefore, that for a target to be exposed, transcription needs

Figure 4. Targeting by MSL1 to the *mof* gene involves DNA sequences embedded in the coding region. MSL1 immunostaining (green) in combination with DNA-FISH (red) for the location of the transgene for insertions containing *mof* with an antisense promoter (A), tubulin-driven antisense *mof* gene (B), endogenous promoter-driven *mof* derivative lacking 670 bp of the 3' end (C), or endogenous promoter-driven 3' end of *mof* (1863–2533 bp) (D). Recruitment of MSL1 to the transgenes is indicated by an arrow. DNA was stained with Hoechst 322 (blue).

to run through the gene whereby, if the signal involves degenerative DNA sequences, the orientation of transcription should not make a difference. Therefore, in order to test this hypothesis we placed a *tubulin* promoter at the 3' end of the *mof* gene such that it would make a *mof* antisense transcript (*tub-mof*^{2.4}AS). If targeting involves a DNA element only to be recognized when the chromatin is in an open conformation, this transgene should retain its binding capacities for MSL1. This construct is, indeed, able to recruit MSL1 (Fig. 4B, top panel; Supplementary Table 1). Interestingly, occasional spreading was observed on one of the lines (<5% of the nuclei) (Fig. 4B, bottom panel). This is possibly caused by the presence of targeting cues toward the 3' end that are now more exposed due to the close proximity of the *tubulin* promoter. In order to test if the 3' end of *mof* is essential for targeting, we created a *mof* construct similar to *mof*^{3.5} but now lacking 600 base pairs (bp) of the 3' end of the *mof* gene (*mof*^{2.9}Δ3prime). This construct, despite being transcriptionally active (Supplementary Fig. 3), failed to recruit MSL1 in all lines observed

(Fig. 4C; Supplementary Table 1), which is in agreement with the overall binding preference of MSLs to the 3' end of genes (Smith et al. 2001; Alekseyenko et al. 2006; Gilfillan et al. 2006). We next tested whether the 3' end of the *mof* gene alone is sufficient to recruit MSL1 in the absence of the surrounding sequences. However, we were unable to observe recruitment of MSL1 on this fragment (Fig. 4D).

Our data suggest that it is most likely that the targeting signal for the MSL complex is a DNA sequence embedded in the coding region of *mof*, only to be exposed when the gene is transcribed, and that the sequences toward the 3' end of *mof* are necessary but not sufficient for MSL1 recruitment as a small DNA element, out of context of the entire gene.

Ectopic recruitment of MSL1 to CG3016 also involves transcriptional activity and sequences toward the 3' end of the gene

In order to test whether transcription-dependent recruitment of the MSL complex was not unique for the *mof*

gene but represented a more general feature of other X-chromosomal genes, we next generated independent transgenes carrying *CG3016* with or without its endogenous promoter sequences (*CG3016*³, *CG3016*^{2.5Δprom}) (Supplementary Table 1). Similar to the *mof* gene, we also observed ectopic recruitment of MSL1 to *CG3016* in the presence of its promoter but not in its absence (Fig. 5A,B). We were next interested in studying the contribution of the sequences toward the 3' end of the *CG3016* gene in MSL1 targeting. For this purpose, we mapped a 339-bp region near the 3' end of *CG3016* that shows maximal MSL enrichment by ChIP and sequence similarity to the previously characterized conserved consensus sequence present on both roX1/2 high-affinity sites (Supplementary Fig. 4; Park et al. 2003). We therefore created transgenes containing 339 bp of this sequence as a monomer [*P*{w⁺ *CG3016*(1581–1920)}]. Interestingly, we observed that similar to the *mof* 3' end, this sequence alone was unable to recruit MSL1 ectopically. Intriguingly however, multimerization [containing five

tandem repeats, *P*{w⁺ *CG3016*(1581–1920) 5mer}] of this sequence restored binding of MSL1, indicating that different from the high-affinity sites that recruit MSLs as an ~300-bp monomer (Kageyama et al. 2001; Park et al. 2003; Oh et al. 2004) but similar to the *mof* situation, the affinity of MSL1 for the *CG3016* sequences is too low to be recognized when placed out of context of the gene itself, but can be revealed upon multimerization (Fig. 5D).

The three previously mapped high-affinity sites (*roX1*, *roX2*, and *18D*) have been shown to maintain their ability to recruit MSL1 in an *msl-3*⁰⁸³ mutant background as a small ~300-bp monomer for roX1/2 and as a multimer in the case of the 18D high-affinity site (Kageyama et al. 2001; Park et al. 2003; Oh et al. 2004). Although *CG3016* is not a high-affinity site, we were interested to test if multimerization of a low-affinity site could overcome the requirement for MSL3. Therefore, to address this issue (as described above in Fig. 2D), we crossed the transgenic flies carrying the *CG3016* multimer with flies expressing MSL2 in an *msl-3*⁰⁸³ mutant background and

Figure 5. Transcription-dependent targeting of MSL1 to *CG3016* and the identification of a targeting sequence toward the 3' end of the gene. MSL1 immunostaining (green) in combination with DNA-FISH (red) on an autosomal *P*-element insertion carrying *CG3016* with (A) or without (B) its endogenous promoter. (C) A 339-bp fragment (represented by a white box) corresponding to nucleotide positions 1581–1920 bp from the transcription start site of *CG3016*. Multimerization of the 339-bp sequence (five tandem repeats) in a wild-type (D) and an *msl-3*⁰⁸³ mutant (E) background. Recruitment of MSL1 to the transgenes is indicated by an arrow. DNA was stained with Hoechst 322 (blue).

analyzed MSL1 binding to female polytene chromosomes. Interestingly, we observed that the *CG3016* multimer failed to recruit MSL1 in an *msl-3⁰⁸³* mutant background (Fig. 5E).

These results show that similar to the *mof* gene, MSL complex recruitment on the *CG3016* gene requires transcriptional activity and sequences embedded within the body of the gene. We also show that even though multimerization of 3' of *CG3016* restores the binding of MSL1 in the wild-type context, the affinity of MSL1 for this *CG3016* sequence is too weak to be maintained in the absence of MSL3.

Discussion

In this study, we provide direct evidence for transcription-dependent recruitment of the MSL complex to X-linked genes. We show that passage of the transcription machinery through the gene is important, whereas the type of promoter and the direction of transcription are not. Our data demonstrate that the recruitment signal lies within the transcribed portion of the gene and that targeting occurs independently of neighboring nucleation sites as we can successfully assess recruitment of X-linked genes on autosomal locations in a transcription-dependent manner. Our data support a model for MSL complex targeting to transcriptionally active X-linked genes that encode a certain combination of small degenerative target sequences in their transcribed region.

It has been proposed previously that one can create MSL complex-binding sites on X-chromosomal regions, normally devoid of the MSL complex, by driving strong transcription via 14 tandem repeats of GAL4-binding sequences (Sass et al. 2003). However, because integration of the EP lines used was random and the sites of insertion were not characterized, the nature of the transcribed unit was unclear, and therefore any assumptions made about the importance of certain sequences, direction, or strength of transcription of the targeted unit remained speculative. Furthermore, since the autosomal EP lines failed to recruit MSL complex, it remained unclear to what extent the X chromosomal context played a role in MSL recruitment on these insertion sites. It may therefore be important to study MSL targeting autonomously, out of the X-chromosomal context, to avoid the complication that may be caused by high local concentration of MSLs on the X chromosome in combination with strong transcriptional activation as previously reported (Sass et al. 2003). Furthermore, in contrast to what has been proposed previously (Sass et al. 2003), our data suggest that transcription alone is unlikely to be the sole targeting signal—first, because many genes on the X chromosome are transcribed but not targeted by MSLs (Alekseyenko et al. 2006; Gilfillan et al. 2006; Legube et al. 2006), and second, as shown in this study, the *mof* gene loses its targeting properties without sequences toward the 3' end while still being transcribed (Fig. 4; Supplementary Fig. 3).

Our data show that most likely the targeting signals for the MSL complex include DNA sequences embedded

in the coding regions of *mof* and *CG3016* only to be exposed when the gene is transcribed. These target sequences are found toward the 3' end and are necessary but not sufficient for MSL1 recruitment, out of context of the entire gene. It remains possible, however, that additional elements also contribute to MSL1 target recognition. Such elements could either be DNA sequences or simply other chromatin-associated factors/modifications that facilitate MSL1 target recognition in a chromatin context. Alternatively, an RNA secondary structure might be involved in MSL recruitment. Such a RNA structure would be a feature of the mRNA sequence. Since the secondary RNA structure formed is independent of the orientation of transcription, the situation created in the case in which *mof* is transcribed in an antisense direction would be similar if not identical to the situation in which *mof* is transcribed in a sense orientation. It is unlikely, however, that a natural antisense transcript is involved in MSL recruitment because a *mof* transgene without a promoter failed to recruit MSL1 where it would still have the potential to produce an antisense RNA molecule (Fig. 1D).

We failed to find any predicted target sequences in the *mof* gene, and although the *CG3016* gene shows some similarity to the *roX1/2* high-affinity consensus sequences, *CG3016* does not behave as a high-affinity site, as judged by MSL3-dependent recruitment. This once again emphasizes the difficulty of predicting MSL target sequences. In agreement with previous studies, we conclude that most probably the nature of these sequences determines the affinity of MSL for its targets and not the level of transcription, as any type of promoter reveals *mof* as a target (Dahlsveen et al. 2006; Gilfillan et al. 2006). Our data may also explain in part the limited predictive power of MSL binding, since targeting only occurs when the gene is transcribed and thus the sequence is exposed. Therefore, MSL-binding predictions should also take into account the transcription state of genes or possibly the chromatin state, for example, by creating a genome-wide DNase I map (Sabo et al. 2004) of the X chromosome.

Unlike the full-length *mof* or *CG3016* genes, the affinity of MSL1 for the 3' end sequences of the *mof* and *CG3016* genes is not sufficient for visualization on polytene chromosomes. Furthermore, even though multimerization of sequences toward the 3' of *CG3016* restores the binding of MSL1 in the wild-type context, the affinity of MSL1 for this *CG3016* sequence is too weak to be maintained in the absence of MSL3. These results show that, on the one hand, there are “high-affinity sites” such as *roX* genes that display transcriptional-independent MSL-targeting sites that are able to recruit MSLs as relatively small DNA sequences (Kageyama et al. 2001; Oh et al. 2004). And, on the other hand, the majority of sites present on the X-chromosome are “low- or moderate-affinity sites” for MSL1 that require exposure of their target sequences presumably by transcriptional activity or artificially by multimerization in order to be recognized as an MSL target sequence. We therefore propose a model of MSL recruitment to the majority of

X-linked target genes based on a combination of active transcription and the presence of DNA target elements.

Future bioinformatics studies encompassing several MSL target genes should reveal whether the 3' ends of X-linked genes harbor a particular consensus sequence, binding sites for other factors, or even a particular secondary structure that together with transcriptional activity contributes to MSL target recognition of X-linked genes.

Materials and methods

ChIP

All ChIP experiments were performed at least three times using independent chromatin preparations as described in Orlando and Paro (1993). Briefly, SL-2 cells were grown in Schneider medium (GIBCO) containing 10% FCS. Cells (1×10^8) were cross-linked with formaldehyde for 8 min. Sonication was performed (26×30 sec) at maximum power (Bioruptor; CosmoBio) in lysis buffer (50 mM HEPES/KOH at pH 7.5, 500 mM NaCl, 1 mM EDTA, 1% Triton X-100, 0.1% DOC, 0.1% SDS + Complete protease inhibitors [Roche]). One-hundred micrograms of chromatin and 3 μ L of polyclonal antibody were used per immunoprecipitation (MSL1, MSL3 raised in rats, and MOF raised in rabbit as described previously; Mendjan et al. 2006). Immunoprecipitated complexes were isolated by adding protein A/G-Sepharose (Roche) followed by four washing steps: twice in lysis buffer, once in DOC buffer (10 mM Tris at pH 8, 0.25 M LiCl, 0.5% NP-40, 0.5% DOC, 1 mM EDTA), and once in TE (pH 8). DNA was eluted in $1 \times$ elution buffer (1% SDS, 0.1 M NaHCO₃) for 20 min at room temperature, followed by reversal of cross-links overnight at 65°C. DNA was purified by a 30-min incubation in 37°C RNase A (0.2 mg/mL) and 2 h in Proteinase K (0.05 mg/mL), followed by phenol/chloroform extraction and DNA precipitation. Each ChIP was resuspended in 100 μ L of TE.

Q-PCR

For the spanning of the cytological location 5c5, primers were designed to space approximately every 500 bp, which is the average size of the chromatin used in this study (primers available on request). The *roX2*, *runt*, and *gapdh* primer pairs are as described in Legube et al. (2006). Q-PCR analysis of the ChIP was performed using the SYBR Green PCR master mix (Applied Biosystem), 100 ng of each primer, and 1 μ L of the immunoprecipitated DNA, in an ABI7500 Real-time PCR Instrument (Applied Biosystem). The formula [percentage ChIP/input] = $[E^{(C_{input}^{ct} - C_{ChIP}^{ct})} \times 100\%]$ (where E represents the primer efficiency) was used to calculate the percentage recovery after ChIP as compared with input. For the analysis of the RNA levels, RNA was first reverse-transcribed using the SuperScript RT (Invitrogen) and 500 ng of random hexamer. One microliter of the cDNA was then subjected to real-time PCR using the SYBR Green PCR master mix (Applied Biosystem) and 10 pmol/ μ L each primer. The primers designed in the middle of the genes in the ChIP experiment were used for the analysis of the transcript levels. For the α -amanitin ChIP experiment, primer pairs were designed to amplify 100- to 200-bp fragments in the beginning, middle, and end of the genes. For the Q-RT-PCR analysis of the transcript levels, the RT reaction was performed as described in the RT-PCR section below. RNA levels were normalized to mitochondrial RNA and depicted as the percentage

recovery of input DNA after α -amanitin treatment compared with mock-treated cells.

RT-PCR

For the RT-PCR in Supplementary Figure 3, third instar larvae were snap-frozen in liquid nitrogen and crushed with a mortar and pestle. RNA was then collected following instructions from the Qiagen RNeasy kit. RT reactions were performed with random hexamers using Invitrogen SuperScriptII. RT reactions were loaded as a dilution series (1, 0.1, and 0.001 μ L of RT mix). For RT-PCR on the transgenes, a primer specific for the CaSpeR vector was used in combination with a *mof*-specific primer. In contrast, for the expression of the endogenous *mof* gene, both primers were encoded within the coding sequence of *mof*.

α -Amanitin treatment

SL-2 cells were grown to a density of $\sim 4 \times 10^6$ and treated for 20 h with either α -amanitin (15 μ g/ μ L; Sigma) or plain insect medium. A total of 1×10^8 cells were used for ChIP, 1×10^6 cells were used to make a nuclear extract as described in Akhtar et al. (2000), and 1×10^6 cells were used for RNA purification. The samples were analyzed by SDS-PAGE followed by Western blot analysis.

Immuno-FISH on polytene staining

Preparation of polytene chromosomes and immuno-FISH were performed as described (<http://www.igh.cnrs.fr/equip/cavalli/Lab%20Protocols/Immunostaining.pdf>). The location of the target genes was detected with specific probes made against the mini-white gene present in our transgenic cassettes. The probes for FISH were generated using random primed dioxigen-dUTP labeling (Roche) of the mini-white gene. MSL1, MSL3, MOF, and MLE antibodies were used at 1:500 dilution (Mendjan et al. 2006). Images were captured with an AxioCamHR CCD camera on a Zeiss Axiovert200M microscope using a 100 $\times$ PlanApocho-mat NA 1.4 oil immersion objective.

Fly genetics

Flies were raised on standard cornmeal-agar-yeast medium at 18°C or 25°C. To generate transgenic flies carrying different insertions, all fragments were cloned using PCR-based strategy (primers available on request) into the p[PCaSpeR]4 vector (Pirrotta 1988), except for *w⁻;P{w⁺ UAS-mof}* and *w⁻;P{w⁺ tub-mof}*, where the *mof* cDNA was cloned downstream from multiple GAL4-binding sites, pP[UAS] (Brand and Perrimon 1993), and downstream from a 2.4-kb *tubulin* promoter sequence, respectively. For the line *y⁻ w⁻;P{w⁺ tub < yellow⁺ > mof}*, the yellow cassette was subcloned from p[P]35 (Struhl and Basler 1993) into the *tub-mof* vector. Transgenic flies were made by *P*-element-mediated transformation of the *w¹¹¹⁸* (also referred to as *w⁻*) recipient stock (Rubin and Spradling 1982). Lines were selected for second or third chromosome insertions. To generate *msl-3⁰⁸³* mutant larvae that carry *P{w⁺ tub-mof}* or *P{w⁺ cg3016 1581–1920 5mer}*, *y⁻ w⁻; msl-3⁰⁸³ [hsp83-*msl-2*]* females were crossed to *w⁻; msl-3⁰⁸³ P{w⁺ tub-mof}/TM6C, Sb Tb* or *P{w⁺ cg3016 1581–1920 5mer}/TM6C, Sb Tb* males. *msl-3⁰⁸³* female larvae, *w⁻; msl-3⁰⁸³ P[hsp83-*msl-2*]/*msl-3⁰⁸³ P{w⁺ tub-mof}* were distinguished by the absence of the *Tb* phenotype. To create *y⁻ w⁻; P{w⁺ tub-mof}* from *y⁻ w⁻; P{w⁺ tub < y⁺ > mof}*, male flies were crossed to female *y w P[70FLP]3F* and progeny was subjected to heat shock at various times during development.*

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Since the stocks were mutant for the endogenous *y* gene, progeny was selected for a loss of the *y*⁺ marker.

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