Domain Requirements of the JIL-1 Tandem Kinase for Histone H3 Serine 10 Phosphorylation and Chromatin Remodeling in Vivo*

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The JIL-1 kinase localizes to Drosophila polytene chromosome interbands and phosphorylates histone H3 at interphase, counteracting histone H3 lysine 9 dimethylation and gene silencing. JIL-1 can be divided into four main domains, including an NH2-terminal domain, two separate kinase domains, and a COOH-terminal domain. In this study, we characterize the domain requirements of the JIL-1 kinase for histone H3 serine 10 (H3S10) phosphorylation and chromatin remodeling in vivo. We show that a JIL-1 construct without the NH2-terminal domain is without H3S10 phosphorylation activity despite the fact that it localizes properly to polytene interband regions and that it contains both kinase domains. JIL-1 is a double kinase, and we demonstrate that both kinase domains of JIL-1 are required to be catalytically active for H3S10 phosphorylation to occur. Furthermore, we provide evidence that JIL-1 is phosphorylated at serine 424 and that this phosphorylation is necessary for JIL-1 H3S10 phosphorylation activity. Thus, these data are compatible with a model where the NH2-terminal domain of JIL-1 is required for chromatin complex interactions that position the kinase domain(s) for catalytic activity in the context of the state of higher order nucleosome packaging and chromatin structure and where catalytic H3S10 phosphorylation activity mediated by the first kinase domain is dependent on autophosphorylation of serine 424 by the second kinase domain. Furthermore, using a lacO repeat tethering system to target mutated JIL-1 constructs with or without catalytic activity, we show that the epigenetic H3S10 phosphorylation mark itself functions as a causative regulator of chromatin structure independently of any structural contributions from the JIL-1 protein.

In Drosophila, histone H3 serine 10 (H3S10)4 phosphorylation by the JIL-1 kinase at interphase functions to maintain active gene expression by serving as a protective epigenetic mark counteracting spreading of H3K9 dimethylation and gene silencing (1–5). Furthermore, JIL-1 is enriched about 2-fold on the male X chromosome and implicated in dosage compensation of transcription due to its association with the male-specific lethal complex (6–9). JIL-1 is a Ser/Thr family tandem kinase that localizes specifically to euchromatic interband regions of polytene chromosomes (6, 10). JIL-1 can be divided into four main domains, including an NH2-terminal domain (NTD), two kinase domains (KDI and KDII), and a COOH-terminal domain (CTD) (6). It has previously been demonstrated that the CTD domain of JIL-1 is sufficient for proper chromatin localization and for rescue of the grossly perturbed polytene chromosome morphology in JIL-1 null mutants (3, 11, 12) and that a JIL-1 construct without the CTD domain has kinase activity for H3S10 despite the fact that it does not localize properly (3, 12). However, in these studies, the properties of the NTD or the relative contributions of the two kinase domains were not determined. Thus, in order to further characterize the domain requirements of the JIL-1 kinase for H3S10 phosphorylation and regulation of chromatin structure, we generated a series of mutated JIL-1 constructs and expressed them in a JIL-1 null mutant background. Our results showed that a JIL-1 construct without the NTD localizes properly to chromatin and rescues mutant polytene chromosome morphology. However, in immunocytochemistry and immunoblot analyses with H3S10ph antibody, no H3S10 phosphorylation could be detected, strongly suggesting that the NTD domain is required for H3S10 kinase activity. Furthermore, we mutated either or both kinase domains to render them “kinase-dead”

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Domain Requirements for JIL-1 Kinase Activity

and assessed their ability to phosphorylate H3S10. The results indicated that both kinase domains are required for kinase activity. In addition, using a LacI/lacO repeat-tethering system (13), we provide evidence that only JIL-1 constructs with H3S10 phosphorylation activity have the capacity to induce a change in higher order chromatin structure from a condensed heterochromatin-like band state to a more open euchromatic interband state.

EXPERIMENTAL PROCEDURES

JIL-1 Fusion Constructs—The full-length LacI-JIL-1 construct (FL) was previously described by Deng et al. (13). LacI-tagged constructs from residues 1–926 lacking the COOH-terminal domain of JIL-1 (ΔCTD), a construct from residues 261–1207 lacking the NH2-terminal domain (ΔNTD), and a COOH-terminal domain construct from residues 927–1207 were cloned into the pUAST vector with the in-frame DNA-binding domain of the LacI repressor from Escherichia coli at the NH2 terminus using standard methods (14). In addition, pUAST constructs were generated using the ‘Transformer™ site-directed mutagenesis kit (Clontech) to introduce K293A (KDI*) and K652A (KDII*) substitutions in the ATP-binding loops for each kinase domain as well as a double mutant combination (KDI*/KDII*). A QuikChange® multisite-directed mutagenesis kit (Agilent) was used to introduce a S425A substitution into the first kinase domain of a full-length JIL-1 cDNA. Subsequently, the mutated JIL-1 cDNA was cloned into the pUASP-HA-FLAG gateway vector according to the manufacturer’s instructions (Invitrogen). For protein purification, a TAP-tagged JIL-1 construct from residues 261–1207 lacking the COOH-terminal domain (TAP-JIL-1) was cloned into the pUAST-NTAP vector (15). The TAP tag consists of calmodulin binding peptide and Protein A. The fidelity of all constructs was verified by sequencing at the Iowa State University sequencing facility.

Drosophila melanogaster Stocks—Fly stocks were maintained at 25 °C according to standard protocols (16). The JIL-1z2 null allele is described by Wang et al. (10) and Zhang et al. (17), whereas the Lac operator insertion line P11.3 and the GFP-LacI fusion line are described by Li et al. (18) and Danzer and Wallrath (19). All transgenic lines were generated by standard P-element transformation (BestGene, Inc.), and expression of the transgenes was driven using hsp70-GAL4/TM6, drivers (19). All transgenic lines were generated by standard P-element transformation (BestGene, Inc.), and expression of the transgenes was driven using hsp70-GAL4 or hsp70-GAL4 drivers introduced by standard genetic crosses. Recombinant JIL-1z2, hsp70-GAL4 chromosomes were generated as described by Li et al. (20). The hsp70 promoter is leaky and was used without heat shock treatment as described previously (12). Expression levels each of the JIL-1 constructs were monitored by immunoblot analysis as described below. All driver lines were obtained from the Bloomington Stock Center. Balancer chromosomes and markers are described by Lindsey and Zimm (21). Viability assays were performed by crossing JIL-transgene/JIL-transgene; JIL-1z2/TM6 flies with JIL-1z2 Hsp70-GAL4/TM6 flies. In these crosses, the TM6 chromosome was identified by the Stubble marker. Consequently, the experimental genotypes could be distinguished from balanced heterozygotic flies by the absence of the Stubble marker. The expected Mendelian ratio of non-Stubble to Stubble flies assuming full rescue by the JIL-1 transgenes in the JIL-1 null background was 1:2 because TM6/TM6 is embryonic lethal. Thus, the percentage of rescue of adult viability for each transgene was calculated as observed non-Stubble flies × 200/observed Stubble flies. In each experiment, the degree of rescue was determined on the basis of the analysis of the genotypes from at least 300 surviving progeny.

Immunohistochemistry—Standard polytene chromosome squash preparations were performed as described by Cai et al. (22) using either 1- or 5-min fixation protocols and labeled with antibody as described by Johansen et al. (23). Primary antibodies used for immunocytochemistry included rabbit anti-H3S10ph (Cell Signaling), rabbit anti-H3K9me2 (Upstate Biotechnology), rabbit anti-JIL-1 (6), chicken anti-JIL-1 (7), and anti-HA antibody (Roche Applied Science). DNA was visualized by staining with Hoechst 33258 (Molecular Probes) in PBS. The appropriate species- and isotype-specific Texas Red-, TRITC-, and FITC-conjugated secondary antibodies (Cappel/ICN, Southern Biotech) were used (1:200 dilution) to visualize primary antibody labeling. The final preparations were mounted in 90% glycerol containing 0.5% n-propyl gallate. The preparations were examined using epifluorescence optics on a Zeiss Axioskop microscope, and images were captured and digitized using a cooled Spot CCD camera. Images were imported into Photoshop, where they were pseudocolored, image-processed, and merged. In some images, non-linear adjustments were made to the channel with Hoechst labeling for optimal visualization of chromosomes.

Immunoblot Analysis—Protein extracts were prepared from adult flies or from dissected third instar larval salivary glands homogenized in a buffer containing 20 mM Tris-HCl, pH 8.0, 150 mM NaCl, 10 mM EDTA, 1 mM EGTA, 0.2% Triton X-100, 0.2% Nonidet P-40, 2 mM Na2VO4, 1 mM phenylmethylsulfonyl fluoride, 1.5 μg/ml aprotinin. Protein separation by SDS-PAGE and electroblot transfer were performed according to standard procedures (14). For these experiments, we used the Bio-Rad Mini PROTEAN III system, electroblotting to 0.2-μm nitrocellulose with transfer buffer containing 20% methanol and in most cases including 0.04% SDS. Primary antibodies for immunoblot analysis included rabbit anti-H3S10ph (Cell Signaling), rabbit anti-histone H3 (Cell Signaling), rabbit anti-JIL-1 (6), chicken anti-JIL-1 (7), and mouse anti-lamin Dm0 (Developmental Studies Hybridoma Bank). The appropriate anti-mouse or anti-rabbit HRP-conjugated secondary antibody (Bio-Rad) (1:3000) was used for visualization of primary antibody. Antibody labeling was visualized using chemiluminescent detection methods (SuperSignal West Pico Chemiluminescent Substrate, Pierce). The immunoblots were digitized using a flatbed scanner (Epson Expression 1680).

Protein Purification and Mass Spectrometry—For TAP-JIL-1 fusion protein purification, the expressed protein was purified from protein extracts of 200 third instar larval salivary glands homogenized in a lysis buffer containing 20 mM Tris-HCl, pH 8.0, 150 mM NaCl, 10 mM EDTA, 1 mM EGTA, 0.2% Triton X-100, 0.2% Nonidet P-40, 2 mM Na2VO4, 1 mM dithiothreitol, 1 mM phenylmethylsulfonyl fluoride, and 1.5 μg/ml aprotinin. Protein lysate was incubated with IgG-agarose beads (Sigma) for 3 h at 4 °C on a rotation wheel. Bound protein beads were
washed with lysis buffer five times. Subsequently, the bound proteins were separated on a 4–12% Tris-glycine gradient gel (Bio-Rad) and stained with Coomassie Blue R250. The TAP-JIL-1 gel band was identified based on size and Coomassie staining intensity and cut out of the gel. The cut band was reduced, alkylated with iodoacetamide, and sent to the University of California, Davis, Proteomics Core Facility for analysis with tandem mass spectrometry using standard methods. Tandem mass spectra were extracted, and charge states were deconvoluted and deisotoped by Scaffold version 3.5.1. All MS/MS samples were analyzed using Sequest version 3.3 (Thermo Fisher Scientific). Sequest was set up to satisfy the principles of parsimony. Using these parameters, a false discovery rate was calculated as 1.7% on the peptide level and 0% on the protein level.

RESULTS

JIL-1 Transgene Expression—In order to further explore the relative contributions of the different JIL-1 domains to H3S10 phosphorylation and chromatin remodeling, we expressed four LacI-tagged JIL-1 UAS P-element insertion constructs constructs transgenically in wild type and JIL-1 null mutant animals. FL, a construct without the COOH-terminal domain (ΔCTD), a construct without the NH2-terminal domain (ΔNTD), and a construct containing only the COOH-terminal domain (CTD) were made (Fig. 1A). A ΔNTD construct has not been previously characterized; however, the LacI-tagged FL, ΔCTD, and CTD constructs used in this study had properties identical to those reported for similar GFP- or CFP-tagged JIL-1 constructs (3, 12) (Figs. 1 and 2). A transgenic line was selected for each construct that expressed at levels comparable with those of endogenous JIL-1 using a hsp-70-GAL4 driver line as illustrated in Fig. 1B. FL rescued all aspects of the JIL-1 null mutant phenotype, including polytene chromosome morphology (Fig. 2B) and viability, and like endogenous JIL-1, FL was up-regulated on the male X chromosome (data not shown). The ΔCTD lacks the COOH-terminal sequences required for proper chromatin localization, leading to mislocalization of the protein (13). However, it does retain its in vivo kinase activity (Fig. 1C), resulting in ectopic H3S10 phosphorylation (12). The ΔCTD rescues autosomal polytene chromosome morphology (Fig. 2B) but only partially rescues that of the male X chromosome in JIL-1 null mutants (12). In contrast, the CTD largely restores JIL-1 null mutant chromosome morphology (Fig. 2B), including that of the male X chromosome (12). Furthermore, when the CTD is expressed in a wild type background, it has a dominant negative effect and displaces endogenous JIL-1 (12).

The NTD Is Required for H3S10 Phosphorylation in Vivo—In order to determine the properties of a JIL-1 construct without the NH2-terminal domain, we expressed the ΔNTD in a JIL-1 null background. A, diagrams of the JIL-1 LacI-tagged constructs analyzed. B, immunoblot labeled with JIL-1 antibody of protein extracts from wild type (wt) and from JIL-1 null salivary glands expressing the FL, the ΔCTD, the ΔNTD, and the CTD, respectively. Labeling with tubulin antibody was used as a loading control. The relative migration of molecular size markers is indicated to the right of the immunoblot in kDa. C, immunoblot labeled with H3S10ph antibody of protein extracts from salivary glands from wild type third instar larvae (wt) and from JIL-1 null salivary glands expressing the FL, the ΔCTD, the ΔNTD, and the CTD, respectively. Labeling with histone H3 antibody was used as a loading control.
null mutant background. As illustrated in Fig. 2B, the ΔNTD restored polytene chromosome morphology to or near the level in wild type preparations (Fig. 2A) and localized correctly to interband regions. Interestingly, in contrast to the ΔCTD, the ΔNTD was not able to phosphorylate H3S10 as documented by immunoblotting of salivary gland protein extracts in Fig. 1C. Furthermore, the polytene squash preparations from larvae expressing the ΔNTD showed that the heterochromatic H3K9me2 mark spread to the chromosome arms (Fig. 2B). Spreading on the X chromosome was especially pronounced in both males and females, as was also the case in the JIL-1 null background (Fig. 2A) and as when the CTD that also is without H3S10 kinase activity (Fig. 1C) was expressed (Fig. 2B). These findings are consistent with the hypothesis that the epigenetic H3S10ph mark itself is necessary for counteracting the spreading of H3K9 dimethylation. Moreover, the results indicate that the NTD of JIL-1 is required for H3S10 phosphorylation to occur in vivo.

H3S10 Phosphorylation Is Sufficient for Chromatin Remodeling—We have previously shown that ectopic targeting of full-length JIL-1 using a LacI-tethering system induces robust H3S10 phosphorylation and a change in higher order chromatin structure from a condensed heterochromatin-like state to a more open euchromatic state (13). However, the surprising finding that the CTD of JIL-1 alone is sufficient to rescue JIL-1 null mutant chromosome defects, including those of the male X chromosome restoring euchromatic interband regions (12) (Fig. 2B), raised the question of whether the CTD also would be sufficient to cause ectopic chromatin remodeling independently of H3S10 phosphorylation. Therefore, we further explored the LacI-tethering paradigm by targeting the various LacI-JIL-1 deletion constructs with and without kinase activity, including the CTD, to chromatin in a condensed heterochromatin-like state. As illustrated in Fig. 3, tethering of the FL or the ΔCTD construct to lacO repeats inserted into the middle of a polytene chromosome band in region 96C1-2

FIGURE 2. Immunocytochemical analysis of polytene chromosome morphology and H3K9 dimethylation in JIL-1 null larvae expressing JIL-1 transgenes. A, comparison of polytene chromosome squash preparations from wild type (wt) and JIL-1 null (z2/z2) third instar larval salivary glands labeled with JIL-1 antibody. B, polytene chromosome squash preparations from JIL-1 null (z2/z2) third instar larval salivary glands expressing the FL, the ΔCTD, the ΔNTD, and the CTD, respectively. Transgenic protein localization (in green) was identified using LacI antibody, H3K9 dimethylation (in red) was identified using H3K9me2 antibody, and DNA (in blue or gray) was labeled by Hoechst.
resulted in ectopic H3S10 phosphorylation and “opening” of the band. In contrast, targeting of GFP-LacI or the CTD resulted in neither H3S10 phosphorylation nor chromatin structure changes. In addition, as illustrated in Fig. 3, the CTD displaces endogenous JIL-1, as reported previously (12), leading to a striking decrease in the levels of H3S10 phosphorylation in the interband polytene chromosome regions. Thus, these findings strongly indicate that the CTD of JIL-1 does not contribute to “opening” of the band when full-length JIL-1 is targeted but rather that these chromatin structure changes are caused solely by H3S10 phosphorylation. Supporting this conclusion, when the ΔNTD was targeted, it accumulated at the target site without any detectable H3S10 phosphorylation or chromatin decondensation and, similarly to the CTD, resulted in decreased overall levels of H3S10 phosphorylation. However, expression of the ΔNTD additionally induced chromatin structure perturbations, especially at the targeting site, as well as ectopic contacts between non-homologous chromatin regions. These results suggest that expression of the ΔNTD, like expression of the CTD, reduced global H3S10 phosphorylation levels by displacing native JIL-1 from the normal binding sites of JIL-1 but that the ΔNTD had an additional dominant negative effect due to the presence of the two kinase domains.

**JIL-1-mediated H3S10 Phosphorylation Requires That Both Kinase Domains Are Catalytically Active**—It has recently been demonstrated that a kinase-dead version of full-length JIL-1 (KDI*/KDII*) in which the crucial lysine for catalytic activity in each of the two kinase domains (Lys293 and Lys652) was changed to alanine, is without kinase activity (13) in a JIL-1 null mutant background. However, these experiments did not address whether one or both kinase domains are functional and required for H3S10 phosphorylation activity. Therefore, we made two additional constructs (KDI* and KDII*), in each of which only one of the lysines was changed to alanine, as diagrammed in Fig. 4A, and expressed them in a JIL-1 mutant background. A transgenic line was selected for each construct that expressed at levels comparable with those of endogenous JIL-1 using an hsp70-GAL4 driver (Fig. 4B). The results showed that the properties of both the single kinase-dead versions, KDI* and KDII*, were indistinguishable from those obtained by expressing a construct where both kinase domains (KDI*/KDII*) were mutated. As illustrated in Fig. 4C, none of the three constructs were able to phosphorylate H3S10, as documented by immunoblotting of salivary gland protein extracts. Furthermore, in polytene squash preparations from JIL-1 null larvae expressing KDI*, KDII*, or KDI*/KDII* (Fig. 5), there was no rescue of chromosome morphology, and spreading of the H3K9me2 mark to the chromosome arms was similar to that observed in JIL-1 null preparations (Fig. 2A). These results strongly indicate that both kinase domains of JIL-1 are required to be catalytically active for H3S10 phosphorylation to occur.

**Phosphorylation of Ser224 Is Required for JIL-1 H3S10 Kinase Activity**—Phylogenetic analysis suggests that JIL-1 is most closely related to the mammalian tandem mitogen and stress-activated kinases 1 and 2 (MSK1/2) in a clade with 95% bootstrap support that is distinct from the RSK (ribosomal S6 p90 kinase) family of tandem kinases (26). The NH2-terminus kinase domain of JIL-1 shares 63% amino acid identity with the corresponding domain in human MSK1 but only 47% identity with that of *Drosophila* RSK (6). MSKs regulate transcription by phosphorylation of transcription factors, including CREB and ATF1 (27), and chromatin-associated proteins, including HMG-14 and histone H3 (28, 29). Similarly to JIL-1, MSK1/2 phosphorylates H3 at Ser10 (29); however, unlike JIL-1 (30, 31), MSK1/2 additionally has histone H3 Ser28 phosphorylation activity (29, 32). The molecular mechanism of MSK activation is complex and requires multisite phosphorylation by upstream kinases and subsequent autophosphorylation of Ser212 and Ser276 by its COOH-terminal kinase domain (33, 34). The JIL-1 kinase is constitutively active and is autophosphorylated (6, 30). Moreover, the region in human MSK1 containing Ser212 in the NH2-terminal kinase domain is highly conserved in JIL-1 (MSK1 Ser212 corresponds to JIL-1 position 424, Fig. 6A), whereas the residue corresponding to MSK1 Ser276 in the linker region has been substituted with a threonine in JIL-1 (JIL-1 Thr280). Thus, to investigate whether these residues also play a role in JIL-1 activity *in vivo*, we expressed a tandem affinity-tagged JIL-1 construct that included the NH2-terminal region.
the two kinase domains, and the linker region (TAP-JIL-1) transgenically in flies. The expressed protein was purified from protein extracts of third instar larval salivary glands and separated using SDS-PAGE, and the enriched TAP-JIL-1 band identified by size and Coomassie Blue staining was cut out and analyzed by tandem mass spectrometry for phosphorylated residues. Of the five peptides recovered containing JIL-1 Thr589, none were phosphorylated; however, in six peptides of 10 containing JIL-1 Ser424, this residue was phosphorylated. Thus, these results indicate that in salivary glands, JIL-1 is phosphorylated at Ser424 but not at detectable levels at Thr589. No other phosphorylated sites were identified in the samples.

In order to determine whether phosphorylation of Ser424 is required for H3S10 kinase activity, we made a JIL-1 construct where Ser424 was mutated to an alanine (JIL-1S424A; Fig. 6B) and expressed it transgenically in a JIL-1 mutant background. A transgenic line was selected that expressed at levels comparable with those of endogenous JIL-1 using a hsp70-GAL4 driver (Fig.

FIGURE 4. Expression of transgenic JIL-1 kinase-dead constructs in a JIL-1 null background. A, diagrams of the JIL-1 V5-tagged constructs analyzed. B, immunoblot labeled with JIL-1 antibody of protein extracts from wild type (wt) and from JIL-1 null salivary glands expressing the KDI*/KDIII*, KDII*, and KDII*, respectively. Labeling with tubulin antibody was used as a loading control. C, immunoblot labeled with H3S10ph antibody of protein extracts from salivary glands from wild type third instar larvae (wt) and from JIL-1 null salivary glands expressing KDI*/KDIII*, KDII*, and KDII*, respectively. Labeling with histone H3 antibody was used as a loading control.

FIGURE 5. Immunocytochemical analysis of polytene chromosome morphology and H3K9 dimethylation in JIL-1 null larvae expressing JIL-1 kinase-dead transgenes. The panels show polytene chromosome squash preparations from JIL-1 null (22/2) third instar larval salivary glands expressing the KDI*/KDIII*, KDII*, KDI*, and FL, respectively. Protein localization (in green) was identified using JIL-1 antibody, H3K9 dimethylation (in red) was identified using H3K9me2 antibody, and DNA (in blue or gray) was labeled by Hoechst.
6D). As illustrated in Fig. 6D, JIL-1*424A had little or no H3S10 phosphorylation activity, as documented by immunoblotting of salivary gland protein extracts. Furthermore, in polytene squash preparations from JIL-1 null larvae expressing JIL-1*424A (Fig. 6C), there was no detectable H3S10 phosphorylation, and spreading of the H3K9me2 mark to the chromosome arms was similar to that observed in JIL-1 null preparations (Fig. 2A). These results strongly indicate that phosphorylation of Ser424 is required for JIL-1 to be catalytically active and for H3S10 phosphorylation to occur. However, it should be noted that although chromosome morphology was still perturbed in JIL-1 null preparations expressing the JIL-1*424A construct, the morphology was discernibly improved compared with preparations expressing KD1*, KDII*, or KD1*/KDII* (Fig. 5).

The properties of the JIL-1 constructs analyzed in the present study, including viability, are summarized in Table 1. The homozygous JIL-1*22 genotype is a late larval lethal with no escapers (17). Interestingly, all of the deletion and mutated constructs with or without H3S10 phosphorylation activity restored some degree of viability, indicating, as suggested previously (12), that multiple and independent mechanisms may be contributing to the lethality of the JIL-1 null mutant.

**DISCUSSION**

In this study, we show that a JIL-1 construct without the NH2-terminal domain is without H3S10 phosphorylation activity despite the fact that it localizes properly to polytene interband regions and that it contains both kinase domains. This is in contrast to a JIL-1 construct without the COOH-terminal domain that, although mislocalized to ectopic chromatin regions, retains its H3S10 phosphorylation capability (12) (this study). Interestingly, although the CTD of JIL-1 binds to the NH2-terminal tail of histone H3, modeling the three-dimensional structure of JIL-1 relative to nucleosome structure suggested that JIL-1 is likely to phosphorylate H3 of one or more nucleosomes some distance away from this binding site.5 Taken together, these findings suggest a model where the NTD of JIL-1 is required for chromatin complex interactions that position the kinase domain(s) for catalytic activity in the context of the state of higher order nucleosome packaging and chromatin structure, whereas sequences in the CTD mediate binding to specific chromatin sites. Furthermore, it has been demonstrated that sequences within both the NTD and CTD are required for enrichment of JIL-1 on the male X chromosome (12).

JIL-1 is a double kinase, raising the question of the function and relative contributions of the two kinase domains. In this study, we show that the properties of both of the single kinase-dead versions of JIL-1, KD1* and KDII*, were indistinguishable from those obtained by expressing a construct where both kinase domains (KD1*/KDII*) were mutated. Neither construct was able to phosphorylate H3S10, rescue JIL-1 null polytene chromosome morphology, or prevent spreading of the H3K9me2 silencing mark, strongly indicating that both kinase domains of JIL-1 are required to be catalytically active for H3S10 phosphorylation to occur. Furthermore, we provide evidence that JIL-1 is phosphorylated at Ser424 in vivo and that this phosphorylation is required for JIL-1 H3S10 phosphorylation activity. This phosphorylation site is conserved in the mammalian MSK family of tandem kinases that are the closest homologs in the databases to the JIL-1 kinase (26). Upon activation of MSKs by upstream kinases, this site has been demonstrated to be autophosphorylated by the COOH-terminal MSK.

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5 W. Cai, unpublished results.
kinase domain and to be required for catalytic activity (33, 34). Thus, the data presented here are compatible with a model where autophosphorylation of Ser424 by the second kinase domain of JIL-1 is similarly required for catalytic activity and H3S10 phosphorylation mediated by the first kinase domain. The molecular mechanism of MSK activation involves multisite phosphorylation that is context- and pathway-dependent (33, 34). In contrast, the JIL-1 kinase is constitutively active (6, 30), and therefore the requirements for its activation are likely to be less complex. Supporting this hypothesis, we did not find evidence for phosphorylation of sites other than Ser424, including Thr589, whose corresponding residue is required to be autophosphorylated for activation of the MSKs.

An important issue of epigenetic chromatin structure regulation by histone phosphorylation is whether the H3S10ph mark itself has the capacity to induce chromatin changes or whether it only plays a reinforcing or maintenance role. Using a LacI tethering system, Deng et al. (13) provided evidence that ectopic H3S10 phosphorylation by the JIL-1 kinase is sufficient to cause striking changes in chromatin packaging from a condensed to an open state. This effect was absent when a kinase-dead LacI-JIL-1 construct that is without H3S10 phosphorylation activity was expressed (13). This indicates that the observed chromatin structure changes depended on JIL-1 kinase-mediated H3S10 phosphorylation and not on the tethering of the LacI-JIL-1 construct itself. However, the surprising finding that the CTD of JIL-1 alone is sufficient to rescue JIL-1 null mutant chromosome defects, including those of the male X chromosome, restoring euchromatic interband regions, suggested that alternative or redundant mechanisms of direct chromatin structure modification may be mediated by the CTD (12). One possibility is that the CTD may serve as a binding platform for an unidentified protein or protein complex that has the ability to induce changes in chromosome morphology but that can function only after proper localization through interactions with the CTD of JIL-1. Another possibility is that binding of the CTD to chromatin by itself is sufficient to alter chromatin structure through bridging interactions with surrounding molecules. However, in this study, we demonstrate using the same experimental paradigm as Deng et al. (13) that LacI-mediated tethering of neither the CTD nor the ∆NTD of JIL-1 resulted in H3S10 phosphorylation or transformation of a polytene chromatin band into an interband and that such chromatin remodeling only is effected by JIL-1 constructs with H3S10 kinase activity. Thus, these findings provide further evidence that the epigenetic H3S10 phosphorylation mark itself functions as a causative regulator of chromatin structure independently of any structural contributions from the JIL-1 protein.


domain of JIL-1 is similarly required for catalytic activity and H3S10 phosphorylation mediated by the first kinase domain. The molecular mechanism of MSK activation involves multi-