Phosphorylation of histone H4 Ser1 regulates sporulation in yeast and is conserved in fly and mouse spermatogenesis

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Sporulation in Saccharomyces cerevisiae is a highly regulated process wherein a diploid cell gives rise to four haploid gametes. In this study we show that histone H4 Ser1 is phosphorylated (H4 S1ph) during sporulation, starting from mid-sporulation and persisting to germination, and is temporally distinct from earlier meiosis-linked H3 S10ph involved in chromosome condensation. A histone H4 S1A substitution mutant forms aberrant spores and has reduced sporulation efficiency. Deletion of sporulation-specific yeast Sps1, a member of the Ste20 family of kinases, nearly abolishes the sporulation-associated H4 S1ph modification. H4 S1ph may promote chromatin compaction, since deletion of SPSt increases accessibility to antibody immunoprecipitation; furthermore, either deletion of Spst or an H4 S1A substitution results in increased DNA volume in nuclei within spores. We find H4 S1ph present during Drosophila melanogaster and mouse spermatogenesis, and similar to yeast, this modification extends late into sperm differentiation relative to H3 S10ph. Thus, H4 S1ph may be an evolutionarily ancient histone modification to mark the genome for gamete-associated packaging.

[Keywords: Saccharomyces cerevisiae; fly and mouse spermatogenesis; genome compaction; histone H4 phosphorylation; kinase; yeast sporulation]

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chromosome condensation in *Tetrahymena* [Wei et al. 1999].

Yeast sporulation is triggered by nutrient deprivation, including nitrogen starvation, growth in nonfermentable carbon sources, or the absence of glucose [Kupiec et al. 1997]. The diploid yeast cell then initiates a transcriptional cascade characterized by expression of several temporally distinct classes of genes: early, middle, mid-late, and late [Mitchell 1994; Chu et al. 1998; Primig et al. 2000]. DNA is replicated, and two meiotic divisions occur during the early and middle stages of the sporulation program. Mid-late and late gene products regulate spore morphogenesis/maturatin [Chua and Herskowitz 1998; Chu et al. 1998]. Haploid spores remain quiescent for an indefinite time until favorable conditions initiate germination. Spores have a specialized wall that imposes volume restrictions on the nuclei and serves as a physical barrier against environmental stress.

Covalent histone modifications have been characterized during early and middle stages of yeast sporulation, temporally coincident with gene regulation, with the occurrence of recombination of double-strand breaks, and with meiotic chromosome condensation. Dynamic H3 and H4 acetylation ([H3ac and H4ac]/deacetylation activates transcription of early and middle genes [Chua and Roeder 1995; Rundlett et al. 1998; Burgess et al. 1999; Choy et al. 2001; Deckert and Struhl 2001]. DNA synthesis occurs during S phase followed by meiosis I and II, which are accompanied by histone phosphorylation [Hsu et al. 2000; Ahn et al. 2005b]. S10ph on both H2B and H3 correlates with meiotic chromosome condensation and disappears during meiotic divisions. Histone ubiquitylation (ub) and methylation (me) are involved in meiotic DNA recombination and middle gene expression [Nislow et al. 1997; Robzyk et al. 2000; Hwang et al. 2003; Wood et al. 2003; Sollier et al. 2004; Yamashita et al. 2004]. It is not known whether histone PTMs influence late gene transcription, or whether they regulate spore-associated genome compaction.

Higher eukaryotic spermatogenesis is conceptually similar to yeast sporulation in the requirement for remodeling and compaction of the genome. Spermiogenesis is marked by two post-meiotic events. In the first event, spermatocytes differentiate into round, and then elongated spermatids. The second major event involves PTMs of histones, replacement of histones by histone variants, then by highly basic transition proteins (TPs), and, finally, by proteasomes [Sassone-Corsi 2002; Govin et al. 2004; Kimmins and Sassone-Corsi 2005]. These changes in histone composition promote efficient genome compaction, sperm function, and increased fertility [Oliva and Dixon 1991; Yu et al. 2000; Cho et al. 2001; Zhao et al. 2001; Meistrich et al. 2003; Lewis et al. 2004]. For example, the timing of H4ac correlates with compaction and may have a direct mechanistic role through binding of the testes-specific bromodomain protein Brdt [Pivot-Pajot et al. 2003]. A large number of phosphorylations occur on the testes-specific histone relatives and replacement proteins, whose molecular mechanisms are not yet elucidated, but may be involved in genome compaction [Oliva and Dixon 1991; Wu et al. 2000; Meetei et al. 2002]. Many components of the spermatogenesis program, including certain histone PTMs, show evolutionary conservation from yeast to mammals. However, it is not known whether histone replacement proteins have a role in flies and yeast, and if not, how lower eukaryotes achieve compaction with only canonical histones.

There is a paucity of information about the role of histone covalent modifications in the broad chromatin-structuring events in gametes. In this study, we describe the occurrence of H4 S1ph during yeast sporulation, as well as during fly and mouse spermatogenesis. H4 S1ph persists after the disappearance of H3 S10ph during gametogenesis in each organism and in yeast is stably present in mature spores. Our findings suggest an evolutionarily conserved role of H4 S1ph during chromatin compaction in the later stages of gametogenesis.

**Results**

*H4 S1ph is observed in sporulating yeast cells and is dependent on Sps1, a middle-sporulation-specific kinase*

Histone H3 phosphorylation occurs during both mitotic and meiotic chromosome condensation [Hsu et al. 2000]. Because H4 S1ph is detected during mitotic chromosome condensation [Barber et al. 2004], we tested whether histone H4 S1 is phosphorylated during sporulation. We used an antibody that specifically recognizes H4 S1ph [Barber et al. 2004; Cheung et al. 2005]. Diploid yeast cells were induced to proceed synchronously through sporulation, and samples were taken at the indicated time points. A strong increase in H4 S1ph occurs at ~10 h in sporulation media [Fig. 1A]. We found that the timing of this modification varies between 8 and 12 h, depending on the severity of starvation, amount of glucose, and aeration of the media [see below].

The timing of the histone H4 phosphorylation appears to coincide with the expression of middle genes. *SMK1* and *SPS1* are Ser/Thr kinases that are expressed in a middle-sporulation-specific fashion and function in parallel pathways to regulate spore wall formation [Friesen et al. 1994; Krisak et al. 1994; Chu and Herskowitz 1998; Iwamoto et al. 2005]. To determine whether these kinases are required for H4 S1ph, we tested diploid strains deleted for either *SPS1* or *SMK1* and carrying plasmid-borne Flag-H4. S1ph in Flag-H4 immunoprecipitates is induced in the smk1Δ strain [Fig. 1B, top panel], but the signal is abolished in the sps1Δ strain [Fig. 1B, bottom panel]. This signal reduction was observed both for Flag-H4 and endogenous H4 [suggesting that mixed octamers immunoprecipitated with Flag antibody]. Thus, sporation-linked H4 S1ph requires Sps1, a middle-sporulation-specific kinase in the Ste20 family of kinases.

*SPS1* encodes a kinase whose transcript levels peak at mid-sporulation and then decrease [Friesen et al. 1994]. We compared the onset of H4 S1ph and induction of Sps1 protein during the sporulation program [Fig. 1C]. Flag-Sps1 expression is induced ~12 h post-induction in
sporulation medium containing 0.05% glucose (where the sporulation program is slightly delayed compared with medium lacking glucose or increased aeration). In induction of H4 S1ph exhibits similar timing as epitope-tagged Sps1 (Barber et al. 2004); however, H4 S1ph persists even after Flag-Sps1 has declined (Fig. 1C). Sps1 is degraded rapidly during purification (data not shown); thus we have been unable to determine whether Sps1 is the direct kinase for H4 S1 during sporulation.

Figure 1. H4 S1ph occurs in sporulation and requires Sps1. (A) Detection of H4 S1ph during sporulation. The wild-type SK1 strain was induced to sporulate, and fractions were collected at the indicated time points. WCEs were prepared, resolved on SDS-PAGE, blotted, and probed either with unmodified H3 [ab1791; Abcam] antibody or with H4 S1ph antibody (Barber et al. 2004). (B) Deletion of Sps1 kinase strongly lowers H4 S1ph, whereas deletion of Smk1 kinase does not affect H4 S1ph. smk1Δ [top panel] and sps1Δ [bottom panel] SK1 strains carrying Flag-H4 were induced to sporulate, and fractions were collected at the indicated time points. Flag-H4 was immunoprecipitated from WCEs, resolved by SDS-PAGE, blotted, and probed with either Flag-HP conjugated antibody (Sigma) or H4 S1ph antibody. SPM* contains 0.05% glucose. (C) Expression of Flag-Sps1 coincides with appearance of H4 S1ph. WCEs were prepared from the sporulating Flag-Sps1 strain, Western-blotted, and probed with Flag and H4 S1ph antibodies.

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Phosphorylation of H3 S10 and H4 S1 are temporally distinct during yeast sporulation

As mentioned above, H3 S10ph occurs during meiotic chromosome condensation, increasing during late prophase and declining as the meiotic divisions occur (Hsu et al. 2000). Because both H3 S10ph and H4 S1ph occur during mitotic chromosome condensation (Barber et al. 2004), it was surprising that S1ph appeared to occur later in sporulation. In fact, H4 S1ph also occurs in response to DNA damage-induced double-strand breaks in vegetative growth (Cheung et al. 2005; Utley et al. 2005), which are similar to meiosis-specific double-strand breaks that occur in meiotic prophase. These observations led us to further examine the timing of H4 S1ph induction during sporulation, especially relative to H3 S10ph and meiotic divisions. Extracts were prepared for Western blotting and, in parallel, cells were DAPI stained to follow the meiotic divisions. The expression of the meiosis-specific Imc2 kinase increases shortly after meiotic induction and increases further as the meiotic divisions are occurring (Fig. 2A, left panel; Benjamin et al. 2003). As previously observed (Hsu et al. 2000), H3 S10ph peaks at ∼4 h coinciding with early sporulation (Fig. 2A) and then declines by 6 h. In contrast, H4 S1ph increases at ∼8 h, when 80% of cells have completed either meiosis I or meiosis II [Fig. 2A]. Thus, while mitotic chromosome condensation is marked with redundant phosphorylation of H3 S10 and H4 S1, these phosphorylations are temporally distinct during sporulation, suggesting separate functions.

We determined the duration of H4 S1ph during sporulation. Western blotting relative to unmodified H3 indicates that it persists to the longest post-meiotic time point tested (30 h) [Fig. 2B]. Thus, unlike transient H3 S10ph, H4 S1ph appears to be stable. We examined S1ph during germination, when the haploid gamete begins to grow in nutrient-rich conditions. H4 S1ph declines between 2 and 4 h following resuspension of the cells in rich YPD medium [Fig. 2B]. Thus, H4 S1ph is a stable modification in spores that is reversed when cells resume mitosis. Induction of H4 S1ph exhibits similar timing as epitope-tagged Sps1 [Fig. 1C]; however, H4 S1ph persists even after Flag-Sps1 has declined [Fig. 1C]. Taken together, these data suggest that a H4 S1ph phosphatase is active during germination but not in spores.

Decreased sporulation in histone H4 S1A substitution mutant

Histone H3 S10ph correlates with mitosis/meiosis; however, substitution mutations in this site do not affect these processes in yeast (Hsu et al. 2000). We investigated the function of H4 S1ph by generating diploid yeast bearing substitution mutations at Ser1 within the sole copy of cellular H4. Cells were collected at the end of the sporulation program, and tetrads were counted under light microscopy to obtain a terminal sporulation frequency. H4 S1A sporulates at approximately one-third of wild-type frequency [Fig. 2C]. We found that H4 R3A sporulates with frequency similar to wild type [Fig. 2C], indicating that the effect of S1A mutation is not likely to be caused by nonspecific alteration of the structure of the H4 tail.
We investigated the localization of H4 S1ph and Sps1

Broad genomic distribution of H4 S1ph and Sps1

We observed by Western analysis that Sps1 expression was transient, whereas H4 S1ph was persistent through sporulation (Figs. 1C, 2B). We next addressed the persistence of H4 S1ph and Sps1 genome association by ChIP. The levels peaked at 10 h in sporulation media and declined thereafter (Fig. 3B). This transience of the ChIP signal is the case both for association at SPS100, DIT1, and other locations (Fig. 3A, right panel). Thus, both H4 S1ph and Sps1 appear to be broadly distributed across the genome during middle sporulation when H4 S1ph and Sps1 are induced.

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Sps1 affects histone ChIP accessibility

To try to reconcile the discrepancy between the Western data (Fig. 2B) and ChIP results (Fig. 3B) with respect to H4 S1ph, we hypothesized that during immunoprecipitation, the H4 S1ph antibody may not be able to access the target histone because of chromatin compaction occurring at later stages of sporulation. To test this, we carried out ChIP using antibodies recognizing unmodified histones H3 and H4, and postulated that the ChIP signals may increase in an sps1Δ strain compared with

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Figure 3. H4 S1ph and Sps1 are broadly distributed on the genome. (A, left panel) Enrichment of H4 S1ph at mid-late and late sporulation genes. ChIP experiments in wild-type SK1 and sps1Δ strains at 0 and 10 h in sporulation medium using an H4 S1ph antibody. The values obtained at SPS100 and DIT1 promoters were normalized to input and are represented relative to the 0 time point set to 1. (Right panel) General recruitment of Flag-Sps1. In addition to formaldehyde, EGS was also used as a cross-linker for the ChIP experiment. ChIP values obtained at 10 h into SPM for TATA regions of DIT1 and PHOS, and the telomeric tip region of the right arm of Chromosome VI are presented as the percentage of input. (B, left and middle panels) H4S1ph levels decrease with time by ChIP. ChIP assays using H4 S1ph antibody in wild-type SK1 and sps1Δ strains at various time points in sporulation medium as shown. The values obtained at the promoter region of SPS100 and telomeric tip region of the right arm of Chromosome VI are presented as the percentage of input. (B, middle panel) Recruitment of Flag-Sps1 in time course at SPS100. ChIP experiments were carried out in wild-type SK1 and in Flag-Sps1 strains at the indicated time points in SPM. The ChIP experiment was done using formaldehyde alone as the cross-linker but reducing the time of Ip from overnight to 4 h. Values obtained at the SPS100 promoter region were normalized to their respective inputs. The normalized values obtained in Flag-Sps1 were converted to fold over the value obtained in the SK1 strain at the identical time point that was set to 1.

DNA volume and nuclear size are increased by mutation of H4 S1A

To more directly investigate chromatin compaction, and to determine whether compaction is due to H4 S1ph and not to other possible effects caused by deleting SPS1, we measured the area of DAPI-stained nucleus in H4 wild-type (Fig. 5A, panels a–d,g) compared with H4 S1A (Fig. 5A, panels d–g) and in smk1Δ (Fig. 5B, panels a–c,g) compared with sps1Δ (Fig. 5B, panels d–g) strains. Sporulating yeast cells were collected, spheroplasted, and adhered to slides. Images of DAPI-stained cells (Fig. 5A [panels a,d], B [panels a,d]) and the corresponding DIC images (Fig. 5A [panels b,c], B [panels b,c]) were digitally captured, merged (Fig. 5A [panels c,f], B [panels c,f]), and the areas of the spore nuclei were digitally measured (Fig. 5A [panel g], B [panel g]). The data were clustered into 0.1-µm² intervals. Larger DAPI-stained nuclei were observed in spores of H4 S1A (mean = 0.96 ± 0.0498) compared with H4 wild type (mean = 0.7838 ± 0.032), and in spores of sps1Δ (mean = 0.7682 ± 0.05) compared with smk1Δ (mean = 0.6458 ± 0.0259) (Fig. 5). The histone substitution strains were treated with proteinase K and RNase A, which likely accounts for their larger nuclei. Thus, absence of H4 S1ph either in the sps1Δ strain or in the H4 S1A substitution mutant strain results in increased nuclear volume.

H4 S1ph persists during differentiation of fly spermatids

We investigated whether H4 S1ph and H3 S10ph also occur during male meiosis and spermatogenesis in Drosophila (Fuller 1993). In spermatocytes undergoing the first meiotic division, metaphase I chromosomes stained strongly against both H4 S1ph [Fig. 6A] and H3 S10ph [Fig. 6A’] antibodies. The H4 S1ph signal was detected on meiotic chromosomes from prophase through telophase for both meiosis I and meiosis II cells (data not shown). In contrast, H3 S10ph staining was only prominent in metaphase and decreased substantially in anaphase and...
Histone H4 Ser1 phosphorylation during gametogenesis

We monitored histone H4 S1ph during mammalian spermatogenesis (Sassone-Corsi 2002). Dissociated mouse cells undergoing spermatogenesis were fractionated using a BSA gradient. Extracts from pooled fractions enriched in spermatocytes, round spermatids, and round elongating spermatids were obtained to examine H4 S1ph and H3 S10ph by immunofluorescence and Western blotting. Consistent with our observations in flies, we detected H4 S1ph by immunofluorescence in spermatocytes and in further differentiated round spermatids, whereas H3 S10ph was observed only in spermatocytes [Fig. 7A, panels a-l]. Again, in agreement with the fly data, H4 S1ph disappears in elongating spermatids (Fig. 7A, panel c).

Western blot analysis was carried out on extracts prepared from the different cell types and probed with H4 S1ph and H3 S10ph antibodies [Fig. 7B]. Again, H3 S10ph dramatically decreases after meiosis, while H4 S1ph persists in post-meiotic stages [Fig. 7B]. The quality of the fractionation procedure is shown by the pattern of H3 S10ph specifically in cells containing condensed meiotic chromosomes (Fig. 7, cf A [panels g-i] and 7B). As an additional control, the same blot was probed with an antibody that detects transition protein 2 (TP2), and as expected, TP2 accumulated in elongating cells [Fig. 7B]. Interestingly, the pattern of TP2 accumulation appeared to correlate with reduction of H4 S1ph [Fig. 7B, cf. lanes 5 and 6].

Discussion

Histones and histone replacement proteins have long been known to be phosphorylated during vertebrate spermatogenesis (Sung and Dixon 1970). Indeed, H4 S1 phosphorylation was discovered in 1970 (Sung and Dixon 1970). However, the significance of this phosphorylation during gametogenesis has been unclear. In this study, we examined S1ph in H4 during gametogenesis and found that the mark is conserved through evolution, appearing in diverse eukaryotes ranging from yeast to the metazoans Drosophila and mouse. In these three organisms, H4 S1ph persists relatively late in the process of gametogenesis compared with meiosis-correlated H3 S10ph. We find that H4 S1ph is induced in yeast nearing completion of meiotic divisions, high levels are maintained to the telophase. The H3 S10ph signal was also not detected in prophase spermatocytes [Fig. 6f, arrowhead]. The antibodies did not cross-react between modified H4 and modified H3 in this assay, as staining of metaphase chromosomes with H4 S1ph antibody was blocked by competition with H4 S1ph peptide, but not by unmodified H4 peptide or H3 S10ph peptide [Fig. 6D–F]. Likewise, staining of metaphase chromosomes with H3 S10ph antibody was blocked by competition with the cognate peptide, but not by unmodified H3 peptide or H4 S1ph peptide [Fig. 6G–I].

Immunofluorescence staining of Drosophila male germ cells undergoing spermatid differentiation revealed certain parallels between differentiation of yeast spores and male gametes in the behavior of H4 S1ph and H3 S10ph. During Drosophila spermatogenesis, as in yeast sporulation, H4 S1ph persisted until late in the terminal differentiation stages, while H3 S10ph levels were strongly reduced by the time meiosis was completed. Thus, in round and early elongating haploid spermatids, nuclei stained brightly for H4 S1ph [Fig. 6B], while staining for H3 S10ph was nearly undetectable in the same cells [Fig. 6B']. The H4 S1ph epitopes persisted in spermatid nuclei as cells grew flagella and elongated and were still detected in nuclei undergoing chromatin compaction and nuclear shaping [Fig. 6C]. Staining with H4 S1ph antibody gradually diminished at the later stages of nuclear elongation and shaping (cf. * vs. brackets in Fig. 6C,C') and was not detected in mature spermatids awaiting individualization. Staining for H3 S10ph was not detected in elongating spermatid nuclei at any stage [Fig. 6C']. As mentioned above, incubation with competing peptides confirmed that the staining in round and elongating spermatid nuclei was due to H4 S1ph [Fig. 6J–M'].
end of sporulation, and the mark is strongly reduced following resumption of growth. We find that a single substitution mutation of H4 S1A lowers sporulation efficiency compared with wild type and increases the size of the nucleus. Thus, this histone mark appears to have a significant role during sporulation.

We do not yet know whether Sps1 is the direct kinase that phosphorylates H4 S1 because we cannot detect in vitro activity of recombinant Sps1 prepared in bacteria, and we have been unable to prepare Sps1 from sporulating yeast (T. Krishnamoorthy, unpubl.), which may be due to assembling spore walls (Lynn and Magee 1970) and abundant proteases (Klar and Halvorson 1975) present in sporulating cells. However, Sps1 is required for H4 S1ph during sporulation (Fig. 1B). Sps1 and H4 S1ph are induced coincidently during mid-sporulation and become chromatin-associated, comparable in both timing and broad locations (Figs. 1C, 3). Deletion of SPS1 or substitution of Ser1 results in an increased volume of DNA and increased nuclear size (Fig. 5). Interestingly, Sps1 is a member of the Ste20/p21-activated kinase family, and yeast Ste20 and its mammalian counterpart, Mst1, have been identified as histone (H2B) kinases during apoptosis (Cheung et al. 2003; Ahn et al. 2005a). H4 S1ph in yeast may be involved in chromatin compaction

H3 S10ph correlates with mitotic and meiotic chromosome condensation in all eukaryotes examined (Gurley et al. 1973; Paulson and Taylor 1982, Hendzel et al. 1997; Cobb et al. 1999; Hsu et al. 2000). However, it is unclear whether H3 S10ph contributes directly to condensation in yeast, because substitution mutation of the site does not alter either mitosis or meiosis (Hsu et al. 2000). Indeed, many histone phosphorylation sites in addition to H3 S10 are detected during mitosis (Goto et al. 1999, 2002, Preuss et al. 2003; Barber et al. 2004; Polioudaki et
al. 2004), but none of these, or combinations of these, have yet been shown to alter growth in yeast (Hsu et al. 2000; W.S. Lo and S.L. Berger, unpubl.). In contrast, our results show reduction of overall sporulation efficiency by single substitution mutation of the H4 S1 site (Fig. 2C).

Our data suggest that S1ph plays a role in genome compaction. H4 S1ph is persistent in spores and is strongly reduced between 2 and 4 h post-germination (Fig. 2B). Consistent with a possible role in broad genome compaction, we detect the mark at many genomic locations throughout the genome (Figs. 3, 4A). Deletion of Sps1 was previously shown to alter transcription of mid-late and late sporulation genes (Friesen et al. 1994). However, because we find both the H4 S1ph mark and the putative kinase Sps1 at many genomic positions by
ChIP, and not specifically at mid-late and late sporulation genes, it appears that the role of H4 S1ph is unlikely to be solely related to induction of these sets of genes. Most directly implicating a role for H4 S1ph in DNA compaction, we observe an increase in the size of the nucleus both in the absence of Sps1 as well as in the H4 S1A substitution mutant (Fig. 5).

H4 S1ph may have a direct role in chromatin compaction within yeast spores. The N-terminal tails of histones are involved in chromatin fiber folding (Moore and Ausio 1997; Carruthers and Hansen 2000), and the X-ray crystal structure of the yeast mononucleosome reveals crystal packing contacts between the H4 tail and the DNA of a neighboring particle, suggesting a possible role in intermonomer contact in vivo (Suto et al. 2000; Zheng and Hayes 2003). In vitro biochemical analysis indicates that amino acids 14–19 of the H4 N-terminal tail, and not tails of the other three core histones, are critical for chromatin folding (Dorigo et al. 2003), and H4 K16ac inhibits the formation of higher-order chromatin structures (Shogren-Knaak et al. 2006). H4 S1ph may oppose H4 K16 acetylation (Utley et al. 2005).

In metazoans, H4 S1ph persists beyond the end of meiosis

Similarities in the processes of sporulation and spermatogenesis, particularly in the drastic reduction in nuclear volume in both processes, led us to examine whether H4 S1ph might also correlate with the timing of chromatin compaction during the latter process. We detect H4 S1ph extending well beyond the time that meiosis-associated H3 S10ph is reduced during Drosophila and mouse spermatogenesis (Figs. 6, 7). In these metazoans, H3 S10ph and H4 S1ph both occur during meiotic divisions and thus may play a role in chromosome condensation. However, following meiotic divisions, the H3 S10ph is dramatically lowered, while H4 S1ph persists during the early stages of the developing spermatids when the genome begins to be compacted (Fuller 1993). This is true in mouse, as H4 S1ph continues to be present beyond the meiotic divisions and begins to be reduced contemporaneously with replacement of histones by the highly basic transition proteins (Fig. 7B). Although we do not yet know whether H4 S1ph has a role in genome compaction in metazoans, our data indicate that H4 S1ph has an additional role beyond meiotic divisions, as is the case in yeast. One clear difference between H4 S1ph in lower eukaryotes compared with metazoans is the persistence of the mark in mature spores and elimination only after germination. Thus, while H4 S1ph may directly promote stable chromatin compaction in mature spores, its role in metazoans may help to compact the genome connected to histone replacement by basic transition proteins (Govin et al. 2004).

While many histone modifications have been correlated with broad genomic mechanisms such as transcription and DNA repair, the role of only a few modifications has been elucidated in higher-level biological processes. Here we report that a central biological process, gametogenesis, is critically controlled in yeast by a single histone modification. The similarities we observe between yeast and metazoans in the persistence of H4 S1ph after the decline of H3 S10ph emphasizes its importance.

Materials and methods

Plasmids and yeast strains

Site-directed mutagenesis was used to generate histone H4 mutations in plasmid pRM204 (HHF2-HHT2 CEN-ARS1 TRP).
Plasmids pTKS4 [hh2 SIA-HHT2 CEN-ARS1 TRP] and pTKS5 [hh2 R3A-HHT2 CEN-ARS1 TRP] were plasmid-shuffled into strain FY1716 [MATα his3Δ200 leu2Δ1 ura3-52 trp1Δ63 lys2-128Δ1 (hh1-h1/Δ1::LEU2 (hh2-h2/Δ1::HIS3 pDM9[HHT1-HHF1 CEN-ARS1 URA3]) to yield strains TY97 and TY98, respectively. Yeast strain ySL151, isogenic to FY1716 but carrying pmR204 instead of pDM9, was used as the wild-type control for histone H4. All the above strains were later transformed with a yCP50 plasmid containing HO to switch the mating type and promote mating to give rise to diploids. The HO plasmid was later removed, and the diploid strains thus generated were TY128 (wt H4), TY129 (H4 S1A), and TY130 (H4 R3A). These strains were used for assessing sporulation frequency.

To study H4 S1p by Western analysis and ChIP, the following strains with an SK1 background were used. Diploid wild-type SK1 [LYN150: MATα/a leu2::hisG/leu2::hisG trp1::hisG lys2-SK1/lys2-SK1 his4-D::NIT1::nmt1 111/a his4-N/111/a ura3-SK1/111/a skp1::LYS2/111/a::LYT2; skp1::111/a strain (MDPY10, genotype identical to LNY150 except having an additional disruption, Δskp1::TRP1 Δskp1::111/a strain (MDPY10, genotype identical to LNY150 except having an additional disruption, Δskp1::LEU2 Δskp1::111/a strain, Myc-Sum1 [DBY36, genotype identical to LNY150 except having ND10/HA::URA3/ND10/HA::URA3 SUMI-MYC::TRP1/SUMI-MYC::TRP1, and FL-SPS1:: URA3 strain. The diploid TY156 strain carrying FL-SPS1 was created by first C-terminal-Flag [2×] tagging the chromosomal SPS1 in a haploid strain with SK1 background by using a TRP marker and following the method as described [Longtine et al. 1998], and the tagging was checked by standard procedures. The haploid Flag-SPS1 strain was thereafter transformed with a HO plasmid as described above to make diploids. Plasmid pF31 [FL-HHT2-HHT2 2p URA3] was used to transform sps1Δ or smk1Δ for Flag immunoprecipitations of histone H4. For monitoring expression of Myc-lme2, strain KSY187 [MATα/a IME2-15XMYC::TRP1/IME2-15XMYC::TRP1 ura3/a ura3 leu2::hisG/leu2::hisG trp1::ΔFA::hisG/ trp1::ΔFA::hisG/lys2::hisG lys2::hisG his3-11, his3-11, 15] [Benjamin et al. 2003] was used.

Culture conditions

A single colony from a fresh plate was used to inoculate 5 mL of YPD/selective SC media for an overnight growth at 30°C. The cells of the overnight culture were harvested, washed with water, and used to inoculate 50 mL of YPA at 0.02 OD600 nm and allowed to grow to 0.5 OD600 nm at 30°C. The YPA culture was allowed to grow to 0.5 OD600 nm at 30°C. The YPA culture was allowed to grow to 0.5 OD600 nm at 30°C. The cells were washed, and finally used to inoculate SPM without glutathione after growth to 0.5 OD600 nm at 30°C. The cells were washed, and finally used to inoculate SPM without glutathione after growth to 0.5 OD600 nm at 30°C. Cells were harvested, resuspended in 1 mL of 4% Formaldehyde for 10 min. The cells were washed three times with 1 mL of H2O and resuspended in 50–100 μL of H2O. Five microliters of the cell suspension was placed on a glass slide to observe the cells. This method was used for monitoring the frequency of sporulation. In the case of measuring the DAPI-stained nuclei, the following procedure was used. Yeast cells were processed as described [Guacci et al. 1994] with the following modifications. Sporulating yeast cells were diluted to 0.1 OD600 nm and fixed with formaldehyde. The extent of spheroplasting was determined by monitoring the cells under the microscope. The concentration of proteinase K was 10 μg/mL. The cells were stained with DAPI, and the images were captured using a fluorescent upright microscope.

Yeast WCE preparation

Yeast cells were harvested by spinning down cultures at ~3000 rpm for 5 min at 4°C. The cells were washed with ice-cold water once and broken in lysis buffer [50 mM Tris-HCl at pH 7.4, 150 mM NaCl, 0.1% NP-40; 5 mM MgCl2; 10 mM NaF; 50 mM β-glycerophosphate; protease inhibitors that included pepstatin, leupeptin, aprotinin, protease inhibitor tablets from Roche and 1 mM PMSF; phosphatase inhibitor cocktail from Sigma) using glass beads and the bead beater (Bio-Spec). The lysate was spun at 14,000 rpm at 4°C for 15 min, and the supernatant/WCE was collected and stored at −80°C. Protein estimation was done using the Bradford protein estimation kit.

Flag immunoprecipitation

Fifty microliters of 50% slurry of Flag M2 agarose resin [Sigma] equilibrated in lysis buffer was used for immunoprecipitation from 1 mg of WCE. The binding reactions were incubated overnight at 4°C on a roller and washed four times with 1 mL of lysis buffer; bound proteins were extracted with 30 μL of 2× sample buffer and heated, and an aliquot was used for checking the expression of Flag fusion proteins. Samples were resolved on 10%–20% SDS-PAGE gel and transferred to a nitrocellulose membrane and probed with the antibody.

ChIP

Fifty-milliliter aliquots of the sporulating cells were collected at various time points, cross-linked with formaldehyde, and processed for ChIP as described [Henry et al. 2003]. For Flag-Sps1 ChIP, cells were first cross-linked with 1.5 mM EGS (100 mM stock prepared in DMSO) for 25 min before cross-linking with formaldehyde for 15 min. The antibodies used for ChIP were α-H4S1ph, α-Flag [M2, Sigma], and H3 [α1791, Abcam]. Primer pairs spanning the TATA regions of SPS100, DIT1, PHOS5, MSE regions of SMK1 and DIT1, and the Telomeric Tip regions of the right arm of Chromosome VI were used to amplify the inputs and eluates obtained by ChIP in a Thermal Cycler [ABI Prism 7000 from Applied Biosystems]. Each amplification was done in duplicate, and the average signal from the eluate was divided by its average input signal for normalizations. The sequence of the primer pairs will be provided on request.

Immunofluorescence on fly testes

Drosophila testes from flies carrying a P[w+] [Ubi-Tubulin] transgene (obtained from C. Gonzales, Barcelona, Spain) were stained for immunofluorescence as in Hime et al. [1996] and Yamashita et al. [2003]. Testes were dissected in 1× PBS and transferred to a siliconized slide; the slide was then frozen in liquid nitrogen, and the coverslip was removed with a razorblade. Samples were then incubated at −20°C in 100% ethanol for 10 min, fixed in 4% formaldehyde in 1× PBS buffer for 7 min at room temperature, permeabilized by two rounds of incubation in 1× PBS, 0.3% Triton X-100, and 0.3% sodium deoxycholate for 15 min each, rinsed with 1× PBST [PBS, 0.1% Triton X-100] for 10 min, then incubated in primary antibody in 1× PBS and 1% BSA overnight at 4°C. H4 S1ph antibody was used at 1:2000 dilution. H3 S1ph antibody [Upstate Biotechnology #RR002] was used at 1:100 dilution. Samples were rinsed...
three times in 1× PBST (1× PBS, 0.1% Triton X-100), then incubated for 2 h at room temperature in Alexa fluor-conjugated anti-rabbit and anti-mouse secondary antibodies diluted in 1× PBS and 3% BSA to concentrations recommended by the manufacturer [Molecular Probes, Inc.]. Samples were rinsed three times in 1× PBST, then mounted in VECTORSHIELD medium with DAPI staining [Vector Lab, Inc. #H-1200]. For peptide competition assays, each primary antibody was incubated with the designated peptide at a final concentration of 10 ng/mL peptide at room temperature for 4 h prior to application to the sample. Images were taken on a Leica TCS SP2 AOBS confocal light microscope and processed in Adobe Photoshop.

Mouse spermatogenic cell fractionation and Western analysis

Spermatogenic cells were fractioned according to the Bellvé method [Bellvé 1998]. A detailed protocol has been published recently [Pivot-Pajot et al. 2003]. Briefly, testes were dissected and processed to obtain a homogeneous cell suspension. The cell suspension was laid on the top of a 2%–4% BSA gradient in an airtight sedimentation unit. Cells were allowed to sediment by gravity for 70 min at 4°C. Then the gradient was fractionated, and cells in each fraction were staged using a phase contrast microscope. Chosen fractions were pooled to obtain cell populations enriched (>80%) in defined pooled stages (spermatocytes, round spermatids, and a mix of round, elongated, and condensed spermatids). Cell pellets obtained after fractionation were disrupted in 8 M urea and quantified by Bradford (Bio-Rad Protein Assay), and Western blots were performed according to standard procedures. Antibodies were used at the following dilutions: H4 S1ph, 1/5000; H3 S10ph [Upstate Biotechnology #06-570], 1/1000; and TP2 (gift from W.S. Kistler, University of South Carolina, Columbia, SC), 1/1000.

Immunofluorescence on mouse tests

A fresh testis was frozen in liquid nitrogen and then applied on a Super frost slide. Fixation was performed in 70% ethanol for 5 min. Slides were dried and stored until use. After a 1× PBST washing, cells were permeabilized in 0.5% saponin, 0.25% Triton, and 1× PBS for ~15 min (twice for 7 min). Blocking was performed in 5% milk, 0.2% Tween, and 1× PBS for 30 min at room temperature. All the following steps (antibody dilution and washes) were performed using the same buffer (1% milk, 0.2% Tween, 1× PBS). H4 S1 phos and H3 S10 phos antibodies were diluted at 1/500 and 1/100, respectively, and the incubations were overnight at 4°C. Slides were then washed three times for 5 min, and an anti-rabbit antibody cross-linked with Alexa 488 [Molecular Probes] diluted at 1/500 was applied to the slides for 30 min at 37°C.Slides were washed and mounted in Mowiol medium with 250 ng/mL DAPI. Image acquisitions were made on an Axiophot [Zeiss] microscope using a -40°C chilled CDD camera [Hamamatsu]. Images were then processed using Adobe Photoshop.

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