Histone H3 threonine 11 phosphorylation is catalyzed directly by the meiosis-specific kinase Mek1 and provides a molecular readout of Mek1 activity *in vivo*

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ABSTRACT

Saccharomyces cerevisiae Mek1 is a CHK2/Rad53-family kinase that regulates meiotic recombination and progression upon its activation in response to DNA double-strand breaks (DSBs). The full catalog of direct Mek1 phosphorylation targets remains unknown. Here, we show that phosphorylation of histone H3 on threonine 11 (H3 T11ph) is induced by meiotic DSBs in *S. cerevisiae* and *Schizosaccharomyces pombe*. Molecular genetic experiments in *S. cerevisiae* confirmed that Mek1 is required for H3 T11ph and revealed that phosphorylation is rapidly reversed when Mek1 kinase is no longer active. Reconstituting histone phosphorylation *in vitro* with recombinant proteins demonstrated that Mek1 directly catalyzes H3 T11ph. Mutating H3 T11 to nonphosphorylatable residues conferred no detectable defects in otherwise unperturbed meiosis, although the mutations modestly reduced spore viability in certain strains where Rad51 is used for strand exchange in place of Dmc1. H3 T11ph is therefore mostly dispensable for Mek1 function. However, H3 T11ph provides an excellent marker of ongoing Mek1 kinase activity *in vivo*. Anti-H3 T11ph chromatin immunoprecipitation followed by deep sequencing demonstrated that H3 T11ph was highly enriched at presumed sites of attachment of chromatin to chromosome axes, gave a more modest signal along chromatin loops, and was present at still lower levels immediately adjacent to DSB hotspots. These localization patterns closely tracked the distribution of Red1 and Hop1, axis proteins required for Mek1 activation. These findings provide insight into the spatial disposition of Mek1 kinase activity and the higher order organization of recombining meiotic chromosomes.

INTRODUCTION

Meiotic recombination initiates with DNA double-strand breaks (DSBs) made by the topoisomerase-like transesterase Spo11 (LAM AND KEENEY 2014). DSBs occur throughout the genome, often but not always in hotspots that in *Saccharomyces cerevisiae* mostly overlap with nucleosome-depleted gene promoters (OHTA *et al.* 1994; WU AND LICHTEN 1994; BAUDAT AND NICOLAS 1997; PAN *et al.* 2011). Repair of meiotic DSBs by recombination helps form physical connections between homologous chromosomes that allow the chromosomes to segregate accurately at the first meiotic division (HUNTER 2015). Because recombination defects can lead to mutations and/or aneuploidy, meiotic DSB repair is highly regulated (SUBRAMANIAN AND HOCHWAGEN 2014; HUNTER 2015).

A critical component of this regulation in yeast is Mre4/Mek1, a meiosis-specific paralog of the Rad53 checkpoint effector kinase (ROCKMILL AND ROEDER 1991; LEEM AND OGAWA 1992). In response to Spo11-generated DSBs, the kinases Tel1 and/or Mec1 (orthologs of mammalian ATM and ATR, respectively) become activated and phosphorylate the chromosome axis-associated protein Hop1 among other substrates (CARBALLO *et al.* 2008; CHENG *et al.* 2013; PENEDOS *et al.* 2015). The FHA (Forkhead-associated) domain of Mek1 then binds phosphorylated Hop1, resulting in Mek1 recruitment to chromosome axes where Mek1 undergoes activation (involving trans-autophosphorylation on T327 in its activation loop) and stabilizes Hop1 phosphorylation via positive feedback (NIU *et al.* 2005; NIU *et al.* 2007; CARBALLO *et al.* 2008; CHUANG *et al.* 2012; PENEDOS *et al.* 2015). Activated Mek1 promotes inter-homolog bias in recombination, that is, the preferential use of a homologous chromosome rather than sister chromatid as the template for DSB repair (NIU *et al.* 2005; CARBALLO *et al.* 2008; GOLDFARB AND LICHTEN 2010; KIM *et al.* 2010; HONG *et al.* 2013; LAO *et al.* 2013; SUBRAMANIAN *et al.* 2016). Mek1 does so in part by phosphorylating the Rad54 protein on threonine 132 (T132) (NIU *et al.* 2007; NIU *et al.* 2009). Rad54 is a member of the Swi2/Snf2 DNA-dependent-ATPase chromatin remodeling family and is a binding partner of the strand exchange protein Rad51 (HEYER *et al.* 2006). Mek1-dependent phosphorylation of Rad54 attenuates the interaction with Rad51, allowing the meiosis-specific strand exchange protein Dmc1 to predominate (NIU *et al.* 2009). Mek1 also directly phosphorylates the T40 residue of Hed1; this stabilizes the Hed1 protein and thereby promotes its function as a negative regulator of Rad51 strand exchange activity (CALLENDER *et al.* 2016). Mek1 also promotes the repair of interhomolog strand invasion intermediates through a pathway required for chromosome synapsis and the generation of crossovers whose distribution shows interference (CHEN *et al.* 2015). Finally, *MEK1* is required for checkpoint arrest or delay of meiotic progression in response to unrepaired DSBs (LYDALL *et al.* 1996; XU *et al.* 1997).

The full array of direct Mek1 phosphorylation substrates remains unknown, as only three direct targets have been definitively proven thus far: Mek1 itself, Rad54, and Hed1 (NIU *et al.* 2007; NIU *et al.* 2009; CALLENDER *et al.* 2016). Additional Mek1-dependent phospho-proteins have been identified by mass spectrometry and other approaches, including T11 of histone H3 (GOVIN *et al.* 2010; SUHANDYNATA *et al.* 2016). However, a number of Mek1-dependent phosphorylation events are known or suspected to be indirect (SUHANDYNATA *et al.* 2016). For example, Mek1 is required for phosphorylation of the synaptonemal complex protein Zip1, but the kinase directly responsible is Cdc7-Dbf4, not Mek1 (CHEN *et al.* 2015). Moreover, H3 T11 phosphorylation has been reported as being catalyzed in vegetative cells by other kinases [the pyruvate kinases Pyk1 and, to a lesser extent, Pyk2 (LI *et al.* 2015)], which could in principle be

regulated by Mek1 in meiosis. Therefore, whether H3 T11 is a direct substrate for Mek1 remains to be established.

Mek1 activity plays out in the context of elaborate higher order chromosome structures. Early in meiotic prophase, sister chromatids form co-oriented arrays of DNA loops that are anchored along a linear proteinaceous axis (ZICKLER AND KLECKNER 1999; KLECKNER 2006). Prominent components of these axes include sister chromatid cohesion proteins (including the meiosis-specific Rec8 subunit), Mek1, Hop1, and another meiosis-specific chromosome structural protein, Red1 (SMITH AND ROEDER 1997; BAILIS AND ROEDER 1998; KLEIN *et al.* 1999; PANIZZA *et al.* 2011).

In cytological experiments, immunostaining foci of recombination proteins are axis-associated, indicating that recombination occurs in proximity to axes (reviewed in ZICKLER AND KLECKNER 2015). However, there is an anticorrelation between the DNA sequences preferentially bound by axis proteins (Rec8, Hop1, Red1) and the DNA sequences that often experience Spo11-induced DSBs, which suggests that recombination usually involves the DNA in chromatin loops rather than the DNA embedded in axes (GERTON *et al.* 2000; BLAT *et al.* 2002; PAN *et al.* 2011; PANIZZA *et al.* 2011). To reconcile this paradox, the "tethered-loop/axis complex" (TLAC) model proposes that DNA segments residing on chromatin loops incur DSBs but are recruited, or tethered, to axes by interactions between recombination proteins and axis proteins (KLECKNER 2006; PANIZZA *et al.* 2011). The TLAC model provides a framework for understanding spatial organization of recombining chromosomes, but there is as yet little direct molecular data demonstrating the proposed functional interactions between axes and DSB sites.

How Mek1 fits into this proposed organization also remains unknown. Immunocytology suggests that Mek1 protein is localized primarily on axes (BAILIS AND ROEDER 1998; SUBRAMANIAN *et al.* 2016), supported by the dependence of Mek1 activity on axis proteins (NIU *et al.* 2007; CARBALLO *et al.* 2008). However, Mek1 exerts its known recombination-controlling activity (directly or indirectly) at sites of DSBs. The TLAC model can account for Mek1 acting at both places, but where Mek1 kinase activity actually occurs remains unexplored because of a lack of a molecular marker for the active kinase.

In this study we demonstrate that Mek1 directly phosphorylates histone H3 T11 in response to meiotic DSBs in *S. cerevisiae*. H3 T11ph is dispensable for Mek1 function during unperturbed meiosis, so the purpose of this phosphorylation remains unclear. Nevertheless, we demonstrate the utility of H3 T11ph as a direct molecular marker for active Mek1 by examining the genome-wide localization of H3 T11ph. Our findings suggest that Mek1 exerts its activity at axis association sites but also across chromatin loops, i.e., spreading beyond the immediate locations predicted by the TLAC model.

MATERIALS AND METHODS

Strains and histone mutagenesis strategy

S. cerevisiae and *S. pombe* strains are listed in **Supplemental Table S1**. *S. pombe* strains were generously provided by G. Smith, Fred Hutchinson Cancer Research Center. Histone gene deletion strains and plasmids expressing H3 T11 mutants from Govin et al. (2010) were generously provided by S. Berger, University of Pennsylvania. *S. cerevisiae* strains were of the SK1 strain background. Because of concerns about effects of plasmid (in)stability on the ability to score phenotypes of histone mutants and to reliably measure meiotic parameters because of cell-to-cell heterogeneity within a culture (see Results), we opted to avoid plasmid shuffle systems that have been used by others (AHN *et al.* 2005; GOVIN *et al.* 2010). Instead, strategies involving stable integration or gene replacement were employed, as follows.

Histone gene replacements: *S. cerevisiae* histone genes are arranged in divergently oriented pairs expressing either H3 and H4 or H2A and H2B; there are two of each pair, i.e., two copies encoding each histone. The S10A and T11V mutations were introduced into plasmid-borne copies of *HHT1* and *HHT2* by QuikChange site-directed mutagenesis (Agilent Technologies). These mutant alleles were then introduced sequentially into SK1 strain SKY165 by one-step gene replacements using DNA fragments containing ≥270 bp arms of homology. Targeting constructs included selectable drug resistance markers: *kanMX4* ~366 bp downstream of the *HHT1* ORF and *hphMX4* ~250 bp downstream of *HHT2*.

Stable integration of histone gene cassettes*:** A histone cassette integration strategy was employed using pRS305-based plasmids (SIKORSKI AND HIETER 1989) integrated into the *leu2::hisG* locus. Integrations were performed to try to maintain balanced gene dosage for the four core histones. The parental strain for the H2A/H2B/H3/H4 histone cassette integrations was created in a multistep process by first transforming a pRS316-based *URA3* histone cassette covering plasmid containing a single copy of each histone gene (pRK12; *HTA1-HTB1*, *HHT2-HHF2*) into diploid SKY165. Next, the histone gene pairs, *HHT2-HHF2* and *HTA1-HTB1* (which are required for proper meiosis (NORRIS AND OSLEY 1987)), were deleted sequentially and replaced with the *hphMX* and *natMX* markers, respectively. The deletions were confirmed by Southern blot and the strain was sporulated to yield a Ura^+, Nat^R, Hyg^R, *MAT*α haploid. A second *MATa** haploid strain was created by sequentially deleting the other (non-essential) histone gene pairs, *HTA2-HTB2* and *HHT1-HHF1*, which were replaced by the *kanMX* and *natMX* markers, respectively, and confirmed by Southern blot. These two haploids were mated to form a compound heterozygote, then tetrads were dissected and resulting haploids carrying all four histone gene-pair deletions were mated to form a histone integration host strain (SKY2283) with the genotype: *hht1-hhf1*Δ*::kanMX/", hht2-hhf2*Δ*::natMX/", hta1-htb1*Δ*::hphMX/", hta2-htb2*Δ*::natMX/", pRK12[CEN/ARS, URA3, HTA1-HTB1, HHF2-HHT2]*.

A parental strain for the H3/H4 histone cassette integrations was created by dissecting tetrads from the *hht2-hhf2*Δ*::natMX/", pRK12* strain described above prior to deletion of *HTA1-HTB1*. This dissection yielded a Ura^+, Nat^R, *MAT***a** haploid that was crossed with the second haploid strain described above (*hta2-htb2*Δ*::natMX, hht1-hhf1*Δ*::kanMX*). Tetrad dissection yielded *MAT***a** and *MAT*α haploid progeny (SKY3166 and SKY3167, respectively) with the following genotype: *hht1-hhf1*Δ*::kanMX, hhf2-hht2*Δ*::natMX, hta2-htb2*Δ*::natMX, pRK12.*

All histone mutant integration constructs were created by QuikChange site-directed mutagenesis. The first was a H3/H4 replacement using a pRS305-based plasmid (pRK77) containing *LEU2*, *HHT2-HHF2* that was linearized by AflII digestion to target integration to *leu2::hisG* and transformed into haploids SKY3166 and SKY3167. The second was an

H2A/H2B/H3/H4 replacement using a pRS305-based plasmid (pRK24) containing *LEU2*, *HTA1-HTB1* and *HHF2-HHT2* that was linearized by AflII digestion and transformed into diploid SKY2283. In both cases, the core-histone covering plasmid pRK12 was counterselected by growth on 5-fluoroorotic acid (FOA). Colony PCR of Leu^+, Ura^- transformants was used to verify the proper integration into the *leu2::hisG* locus using primer sets flanking both junctions as well as verification of the mutations in *hta1* and *hht2* by engineered restriction enzyme site polymorphisms and/or sequencing. In the case of the SKY3166/3167 transformants, haploid integrants were subsequently mated to create diploids. SKY2283 hemizygous integrants were sporulated to produce haploid progeny that were then mated to create homozygous diploids.

S. cerevisiae and *S. pombe* cultures

S. cerevisiae was cultured at 30°C with asynchronous vegetative (cycling) cultures in YPD (1% yeast extract, 2% peptone, 2% dextrose). Camptothecin treatment (20 µM) was performed for 2 hr at 30° in 250 ml flasks shaking at 250 rpm in 10 ml cultures of SKY165 at an initial cell density of ~9×10^7 cells/ml. An untreated culture was incubated in parallel, while a separate 10 ml aliquot in a vented T-75 flask was exposed to X-rays for 60 min at room temperature using an X-RAD 225C X-ray irradiator (Precision X-ray, Inc.) corresponding to a dose of 400 Gy. Alternatively, 10 ml of culture at ~7×10^7 cells/ml was exposed to X-rays for 60 min on ice, with untreated cells also held on ice. With both exposure conditions, cells were subsequently allowed to recover at 30°, shaking at 225 rpm for 60 min (room temperature exposure) or 30 min (exposure on ice) before fixing in 20% trichloroacetic acid (TCA), pelleting and storage at -80° until extract preparation.

For inhibition of Mek1-as *in vivo*, an SKY3095 culture was divided equally four hours after transfer to sporulation medium and 10 µl 100% DMSO was added to half while the other received 1 µM final concentration of 1-NA-PP1 (1-(1,1-Dimethylethyl)-3-(1-naphthalenyl)-1H-pyrazolo[3,4-d]pyrimidin-4-amine) dissolved in DMSO (WAN *et al.* 2004). The return-to-growth recombination assays using *arg4* heteroalleles were carried out in triplicate as described (MARTINI *et al.* 2006). Pulsed-field gel electrophoresis (PFGE) and Southern blotting on DNA from meiotic cultures prepared using the SPS method was performed as described (MURAKAMI *et al.* 2009). Plasmid shuffling and meiotic cultures using plasmids and the SK1 histone gene deletion strain obtained from S. Berger were carried out as described (GOVIN *et al.* 2010).

S. pombe haploid *pat1-114* sporulation was carried out as described (HYPPA AND SMITH 2009). For *S. cerevisiae* meiotic cultures, strains were thawed on YPG plates (1% yeast extract, 2% peptone, 3% glycerol, 2% agar) and incubated for ~2 days, then streaked for single colonies on YPD plates and grown ~2 days. Single diploid colonies were inoculated in 5 ml YPD and grown overnight. Cultures were diluted in YP+1% potassium acetate presporulation medium to ~1.2×10^6 cells/ml, grown for 13.5 hours at 225 rpm for ChIP and 250 rpm for all other experiments. Cells were pelleted, washed in sterile water and resuspended in the same preculture volume of 2% potassium acetate to a density of ~$2–3 \times 10^7$ cells/ml. This corresponds to 0 hr of the meiotic time course. Sporulation was at 225 rpm for chromatin immunoprecipitation (ChIP) and 250 rpm for all other experiments. Meiotic progression was assessed in culture aliquots fixed with 50% ethanol and stained with 5 µg/ml 4',6-diamidino-2-phenylindole (DAPI).

Whole-cell extracts and western blotting

Culture aliquots of OD_{600} = 10 for *S. pombe* or ~3.2×10^8 cells for *S. cerevisiae* were washed in 20% TCA, pelleted and stored at -80°C until ready for use. Aliquots were thawed,

resuspended in 20% TCA and disrupted by bead beading at 4° using 0.5 mm zirconia/silica or glass beads and monitored microscopically until near complete disruption was observed. Samples were collected by centrifugation, then washed with 5% TCA and the pellet was resuspended in 1× NuPAGE LDS Sample Loading Buffer (Life Technologies Corp.) with 100 mM dithiothreitol (DTT). Samples were separated on 12% bis-Tris NuPAGE gels in 1× MOPS or MES running buffer (Life Technologies Corp.) or 15% Laemmli gels (LAEMMLI 1970). Proteins were blotted to polyvinyldifluoride (PVDF) membranes by semi-dry electrophoretic transfer using the iBlot system (Life Technologies Corp.) or in Tris-glycine (25 mM Tris base, 192 mM glycine, 10% methanol, 0.04% sodium dodecyl sulfate) at 100 mA constant for 70 min (TransBlot SD Transfer Cell, Bio-Rad Laboratories, Inc.). Membranes were air dried, then incubated with one of the following rabbit primary antibodies diluted in 5% non-fat milk (NFM) in Tris-buffered saline-Tween buffer (TBST; 25 mM Tris-HCl pH 7.4, 137 mM NaCl, 2.7 mM KCl, 0.1% Tween 20): anti-H3 polyclonal (Abcam 1791) diluted 1:10,000; anti-H3 T11ph mononclonal (EMD Millipore 05-789) diluted 1:1000; anti-H3 T11ph polyclonal (Active Motif 39151) diluted 1:1000; anti-H3 S10ph monoclonal (EMD Millipore 05-817) diluted 1:1000; anti-H3 S10ph polyclonal (EMD Millipore 06-560) diluted 1:1000; or anti-H2A S129ph polyclonal (Abcam 15083) diluted 1:500. The polyclonal secondary antibody used was horseradish peroxidase-conjugated goat anti-rabbit (Pierce/ThermoFisher Scientific 31462 or 31460) diluted 1:10,000 in TBST with visualization by the ECL-Plus kit (GE Healthcare Ltd.) exposed to chemiluminescent film or charged-coupled device (CCD) camera (Imagestation, Eastman Kodak Company).

Validation of anti-phospho-H3 antibodies

Two commercial anti-H3 T11ph antibodies yielded Spo11-dependent bands at the expected size for H3 on western blots, but the monoclonal gave more robust signal with less background (**Figure 1B**). To more definitively characterize the specificity of these antibodies, we incubated them with synthetic peptide arrays containing different H3 modification states (Active Motif MODified histone peptide array)(**Supplemental Table S2**). The monoclonal anti-H3 T11ph antibody reacted strongly with all peptides containing T11ph regardless of other modifications present, unless S10 was also phosphorylated, in which case reactivity was strongly or completely lost (**Supplemental Figure S1Ai**). This monoclonal antibody was highly specific, as little to no cross-reactivity was observed for unmodified H3 peptides, H3 peptides carrying other modifications, or peptides from other histones, including peptides phosphorylated at other sites (H3 S10ph, H3 S28ph, H4 S1ph, H2A S1ph, H2B S14ph) (**Supplemental Figure S1Ai**). In a more limited analysis, the polyclonal anti-H3 T11ph antibody bound specifically to a peptide with trimethylated H3 K9 (K9me3) as well as T11ph, but not to unmodified or S10ph peptides from H3 or full-length unmodified histones (**Supplemental Figure S1B**). However, this polyclonal antibody showed substantial non-histone cross-reactivity against yeast whole-cell extracts that was not observed for the monoclonal anti-H3 T11ph antibody (**Figure 1B**).

Both the monoclonal and the polyclonal anti-H3 S10ph antibodies we used reacted with phospho-S10 H3 peptide on dot blots, but with some background signal for full-length histone H3 (**Supplemental Figure S1B**). Similarly, the polyclonal anti-H3 S10ph antibody detected S10ph on the peptide array, including in the context of other nearby modifications, unless T11 was also phosphorylated (**Supplemental Figure S1Aii**). Again, however, modest cross-reactivity was seen with other histone H3 and H4 peptides, thus the anti-S10ph antibodies are less specific than the monoclonal anti-T11ph antibody.

***In vitro* kinase assays**

GST-Mek1 and GST-mek1-as were affinity purified on glutathione sepharose as described (NIU *et al.* 2009; LO AND HOLLINGSWORTH 2011).

Radiolabeling method: Reactions included 2 µg of recombinant *S. cerevisiae* histone H3 or 5 µg H3 1-20 peptides, 250 ng GST-Mek1, 0.4 mM ATP and 10 µCi [γ-^{32}P]-ATP (6000 Ci/mmol; PerkinElmer, Inc.) in 25 µl total volume in a buffer containing 50 mM HEPES-NaOH pH 7.5, 150 mM NaCl, 10 mM $MgCl_2$, 0.5 mM DTT and 1× each of Roche phosphatase and protease inhibitor cocktails. Reactions were incubated at 30° for 30 min then resolved on 12% bis-Tris NuPAGE gels in 1× MES running buffer and transferred to PVDF via the iBlot system or Coomassie stained and dried for autoradiography on a Fujifilm FLA 7000. Primary antibody was rabbit anti-H3 T11ph polyclonal (Active Motif 39151) diluted 1:500, with secondary antibody and detection carried out as described above.

Semi-synthetic epitope method: GST-Mek1-as target labeling and detection followed previously described methods (NIU *et al.* 2009; LO AND HOLLINGSWORTH 2011). Reactions included 2 µg of recombinant *S. cerevisiae* histone H3, 2 µg GST-Mek1 or 0.76 µg GST-Mek1-as, 0.4 mM ATPγS or 6-Fu-ATPγS (N^6-furfuryladenosine-5'-O-3-thiotriphosphate, Axxora, LLC), and 0.2 mM ATP in 25 µl total volume in a buffer containing 50 mM Tris-HCl pH 7.5, 150 mM NaCl, 10 mM $MgCl_2$ and 0.5 mM DTT. Reactions were incubated at 30° for 30 min, then *p*-nitrobenzyl mesylate (PNBM in DMSO, Abcam/Epitomics 3700-1) was added to 2.5 mM and incubated at room temperature for 90 min. Samples were electrophoresed on 4–12% bis-Tris NuPAGE gels in 1× MES running buffer, followed by semi-dry transfer to PVDF at 25 V constant for 60 min. Membranes were blocked in 5% NFM-TBST, primary antibodies were rabbit anti-thiophosphate ester monoclonal (Abcam/Epitomics 2686-1) diluted 1:5000 or rabbit anti-H3 T11ph monoclonal (EMD Millipore 05-789) diluted 1:1000, with secondary antibody and detection carried out as described above.

ChIP-sequencing

The ChIP-seq protocol was based on a previously described method (WAL AND PUGH 2012). Two independent wild type (SKY165) and one *spo11-Y135F* (SKY198) meiotic cultures were prepared as described (MURAKAMI AND KEENEY 2014) and 4×10^9 cells were harvested at 3 and 4 hr (wild type), and 3.5 hr (*spo11-Y135F*) after the meiosis induction. Cells were fixed with 1% formaldehyde for 15 min at room temperature, with mixing at 50 rpm. Crosslinking was quenched by adding glycine to 131 mM for 5 min, cells were washed with water, resuspended in ice-cold ST buffer (10 mM Tris-HCl pH 7.4, 100 mM NaCl and 1× each of Roche phosphatase and protease inhibitor cocktails). To compare different samples, we used *S. pombe* cells as a spike-in control. *S. pombe* cells (SKY2594) harvested at 4.5 hr in meiosis were fixed and washed with the same condition described above. An aliquot of 4×10^7 *S. pombe* cells (1% of the number of *S. cerevisiae* cells) were added to each sample.

Cells were resuspended in FA lysis buffer (50 mM HEPES-NaOH pH 7.5, 150 mM NaCl, 2 mM EDTA, 1% Triton X-100, 0.1% sodium deoxycholate, 10 µg/ml each of leupeptin, pepstatin A, and chymostatin, 1 mM PMSF, 1× each of Roche phosphatase and protease inhibitor cocktails) and disrupted using zirconia/silica beads (0.5 mm, Biospec Products, Inc. 11079105z) and a FastPrep-24 (MP Biomedicals) with 8 rounds of shaking at 6.5 m/s for 60 seconds. Lysates were pelleted by centrifugation at 15,000 rpm for 5 min at 4°, washed with NP-S buffer (0.5 mM spermidine, 0.075% IGEPAL CA-630, 50 mM NaCl, 10 mM Tris-HCl pH 7.4,

10 mM $MgCl_2$, 2 mM $CaCl_2$, 10 µg/ml each of leupeptin, pepstatin A, and chymostatin, 1 mM PMSF, 1× each of Roche phosphatase and protease inhibitor cocktails) and resuspended in 3.6 mL NP-S buffer with 1 mM 2-mercaptoethanol. The resuspended pellet (chromatin) was solubilized by digestion with 25 units/ml of micrococcal nuclease (Worthington Biochemical Corp.) at 37° for 20 min. Digestion was terminated by adding EDTA to 10 µM and SDS to 0.05%. Chromatin was further solubilized by sonication (Biorupter Standard, Diagenode) on highest setting for two rounds of 30 sec with a 30 sec intervening rest. Solubilized chromatin was isolated by centrifugation at 16,000 rpm for 10 min at 4°, pooled and divided into two equal volumes.

Twenty µg of each antibody was added to the MNase-treated chromatin samples and incubated at 4° overnight on a rotisserie mixer [Antibodies: rabbit anti-H3 pAb (Abcam 1791); rabbit anti-H3 T11ph mAb (EMD Millipore 04-789)]. Immunoprecipitation was carried out by adding 200 µl protein G Dynabeads (Life Technologies Corp.) and incubating at 4° for 90 min on a rotisserie mixer. Beads were washed with 1 ml of the following buffers: NP-S buffer, FA lysis buffer, 2× FA high salt buffer (FA lysis buffer containing 1 M NaCl), 2× FA wash 2 buffer (FA lysis buffer containing 0.5 M NaCl), 2× FA wash 3 buffer (10 mM Tris-HCl pH 8, 250 mM LiCl, 2 mM EDTA, 1% IGEPAL, 1% sodium deoxycholate) and TE wash buffer (10 mM Tris-HCl pH 8, 1 mM EDTA, 0.5% Triton X-100). Bound nucleosomes were eluted, reverse crosslinked, treated with RNase A and Proteinase K as described (MURAKAMI AND KEENEY 2014). DNA was purified using PCR purification kit (Qiagen) and separated on a 1.5% agarose gel. Mononucleosome-sized DNA (~150 bp) was extracted from the gel and prepared for 50 nt paired-end sequencing on the Hiseq platform (Illumina, Inc.) following standard Illumina protocols. Sequencing was performed at the Integrated Genomics Operation of Memorial Sloan-Kettering Cancer Center.

Paired-end 50 nt reads were mapped to the *S. cerevisiae* reference genome (sacCer2) and the Sanger Center's *S. pombe* genome version of 7 August 2010 using BWA (version 0.7.12-r1039) MEM (LI 2013). Paired reads with an insert size more than 250 bp were filtered out and the rest were converted into coverage maps. All downstream analyses were carried out using R (http://www.r-project.org/) (R DEVELOPMENT CORE TEAM 2012). Each coverage map was normalized per 1000 reads from *S. pombe* chromosomes I and II. Cumulative S1-seq values were generated by calculating the cumulative sum of the top strand (or bottom strand) reads of S1-seq data (MIMITOU *et al.* 2017) from the midpoint of each hotspot to 2 kb downstream (or upstream).

Data availability

Plasmids and strains are available on request. ChIP-seq data are available at the Gene Expression Omnibus (GEO), accession number pending.

RESULTS

H3 T11 phosphorylation during meiosis is a response to DSBs

As part of a larger effort to identify meiotically regulated histone modifications in *S. cerevisiae*, we performed western blots on meiotic whole-cell extracts with antibodies to H3 T11ph. Under these conditions, signal was undetectable in mitotically cycling, premeiotic (G1-arrested, 0 hr), or early meiotic (through 2 hr) cultures, but accumulated transiently during meiosis with a maximum at ~3 to 5 h (**Figure 1Ai, 1B**). This signal diminished as cells completed the first meiotic division (~7 hr; **Figure 1Ai, 1B, 1C**). These findings agreed with studies reported while this work was in progress (GOVIN *et al.* 2010).

The anti-H3 T11ph signal occurred when DSBs are usually maximal under these conditions [~3 to 5 hr (e.g., THACKER *et al.* 2014)], and coincided with an increase in H2A S129 phosphorylation (γ-H2A) (**Figure 1Ai**), which is formed by Mec1 and Tel1 kinases in response to meiotic DSBs (MAHADEVAIAH *et al.* 2001; SHROFF *et al.* 2004). These results suggested H3 T11ph might be a DSB response, but H3 T11ph signal also coincided with an increase in H3 S10 phosphorylation (**Figure 1Ai**), which is DSB-independent (HSU *et al.* 2000).

We therefore examined genetic requirements for H3 T11ph. The modification was undetectable in a strain with catalytically inactive Spo11 (*spo11-Y135F*; **Figure 1Aii, 1B**). As expected, induction of higher γ-H2A signal was not seen in *spo11-Y135F,* but H3 S10ph was induced (**Figure 1Aii**). H3 T11ph appeared in a *rad50S* strain, in which DSBs form but persist with unresected 5′ ends, so DSB resection is dispensable (**Figure 1Aiii**). H3 S10ph was unaffected in this mutant, but elevated γ-H2A levels persisted to late time points consistent with unmitigated Tel1 activity (USUI *et al.* 2001).

H3 T11ph appeared and disappeared in *rad51*Δ with kinetics similar to wild type (**Figure 1Av**), but persisted at high levels in *dmc1*Δ (**Figure 1Avi**) (a different antibody was used for these blots, discussed below). Both *rad51*Δ and *dmc1*Δ have defects in meiotic DSB repair (note the persistent γ-H2A), but with a more complete block in *dmc1*Δ (BISHOP *et al.* 1992; SHINOHARA *et al.* 1992). Meiotic arrest is also nearly complete in *dmc1*Δ, whereas divisions occur in *rad51*Δ after a delay (**Figure 1C**) (BISHOP *et al.* 1992; SHINOHARA *et al.* 1992).

To determine whether this persistent H3 T11ph signal was due to persistent DSBs or to meiotic arrest, we examined an *ndt80*Δ mutant. Ndt80 is a transcription factor needed for pachytene exit (XU *et al.* 1995; CHU AND HERSKOWITZ 1998), and DSB repair defects cause arrest via checkpoint kinase-mediated inhibition of Ndt80 (TUNG *et al.* 2000; GASIOR *et al.* 2001). H3 T11ph did not persist in an *ndt80*Δ mutant and instead peaked at 4 h at a slightly lower level than in wild type (**Figure 1Aiv**). This agrees with a recent report demonstrating H3 T11ph appearance and disappearance by western blotting and immunofluorescence of spread chromosomes (SUBRAMANIAN *et al.* 2016). Therefore, H3 T11ph persistence correlates with continued presence of meiotic DSBs (as in *dmc1*Δ), but not with arrest. This behavior contrasts with that of a different Mek1 substrate, Hed1, which remains phosphorylated in *ndt80*Δ mutants (PRUGAR *et al.* 2017). Both γ-H2A and H3 S10ph persisted at high levels in *ndt80*Δ (**Figure 1Aiv**) (HSU *et al.* 2000), suggesting these modifications require pachytene exit for removal (SUBRAMANIAN *et al.* 2016).

Because the H3 N-terminal tail has many potential modification sites (**Figure 1D**) and a different antibody not used in our studies cross-reacts between H3 T11ph, H3 S10ph and other modifications (NADY *et al.* 2008), we sought to validate the antibody specificity for the anti-H3 T11ph antibodies we used. Both the monoclonal and polyclonal anti-H3 T11ph antibodies were specific but did not detect H3 T11ph if S10 was also phosphorylated (**Supplemental Figure S1**

and Materials and Methods). The monoclonal gave a more robust signal with less background for non-histone proteins (**Figure 1B**), so we used this antibody for most subsequent experiments. Two different anti-H3 S10ph antibodies recognized their cognate modification, but not if T11 was also phosphorylated. These anti-H3 S10ph antibodies showed significant cross-reactivity to other histones and modifications (**Supplemental Figure S1** and Materials and Methods).

To test if DNA lesions could also give rise to elevated T11 phosphorylation during vegetative growth, cells were treated with X-rays or camptothecin. These DNA damaging agents failed to yield a detectable level of H3 T11ph despite inducing DNA damage responses as evidenced by increased γ-H2A (**Figure 1E**). Thus, high levels of H3 T11ph are largely if not exclusively specific to meiosis. The strength of the meiotic H3 T11ph signal as compared to the undetectable levels under these blotting conditions for cycling or premeiotic cells or the *spo11-Y135F* mutant indicates that the amount of H3 T11ph formed in meiosis is vastly greater than what has been reported to be formed by pyruvate kinase during vegetative growth (LI *et al.* 2015).

H3 T11ph in response to DSBs in *S. pombe* meiosis

To determine if meiotic H3 T11ph is evolutionarily conserved, we analyzed synchronous meiosis in *S. pombe* haploid *pat1-114* mutants (BAHLER *et al.* 1991). H3 T11ph appeared transiently at ~4–5 hr after the initiation of meiosis and was not detected in a mutant lacking Rec12 (the Spo11 ortholog) or in vegetative growth (**Figure 2**). H3 T11ph appeared after a Rec12-dependent increase in γ-H2A that started around 3–3.5 hr, when DSBs typically appear under these conditions (CERVANTES *et al.* 2000). (The initial wave of γ-H2A signal at or before 2 hr is Rec12-independent (**Figure 2B**) and possibly associated with DNA replication.) These results indicate that H3 T11ph forms in response to DSBs in *S. pombe*. H3 T11ph appeared and disappeared with apparently normal kinetics in a *rad50S* mutant in contrast to γ-H2A, which persisted at high levels (**Figure 2C**).

H3 S10ph also appeared during meiosis, but unlike in *S. cerevisiae*, this modification occurred later than H3 T11ph (**Figure 2A**). In the *rec12* mutant H3 S10ph was observed earlier than normal and was largely gone by 6 hr (**Figure 2B**). This result is consistent with accelerated meiotic progression in *rec12* mutants (DOLL *et al.* 2008), and indicates that both appearance and disappearance of H3 S10ph are developmentally regulated.

H3 T11 is a direct target of Mek1 kinase

The timing and genetic control of H3 T11ph in *S. cerevisiae* suggested that a meiosis-specific, DSB-responsive kinase was responsible. Mek1 expression coincides with H3 T11ph from 3–7 hr in meiosis (CARBALLO *et al.* 2008), and the T11 sequence context matches the Mek1 target consensus (RXXT; **Figure 1D**) (MOK *et al.* 2010; SUHANDYNATA *et al.* 2016). We therefore treated a *dmc1*Δ strain expressing an ATP-analog sensitive *mek1* allele (*mek1-as*) with an inhibitor specific for the mutated Mek1 kinase, 1-NA-PP1 (WAN *et al.* 2004). Inhibitor addition at 4 hr caused rapid disappearance of H3 T11ph within the first hour (**Figure 3A**). This result demonstrates that Mek1 activity is necessary to maintain H3 T11 phosphorylation, and further implies that this modification is dynamic with a half-life much shorter than one hour.

This result agreed with prior findings demonstrating that H3 T11ph is reduced or absent in a *mek1*Δ mutant (GOVIN *et al.* 2010). However, these findings did not establish whether H3 T11 is a direct target of Mek1. To address this question, we carried out two types of *in vitro* kinase assay using GST-tagged Mek1 purified from meiotic *S. cerevisiae* cells (WAN *et al.* 2004;

NIU *et al.* 2007). First, we used [γ-^{32}P]ATP and full-length H3 or synthetic H3 peptides as substrates (**Figure 3B**). GST-Mek1 was visible in all lanes by Coomassie staining (**Figure 3B**, bottom panel) and its activity was confirmed by its ability to autophosphorylate **(Figure 3B**, top panel) (NIU *et al.* 2009). GST-Mek1 was able to phosphorylate full-length H3 and a peptide representing H3 amino acids 1-20 (**Figure 3B**, top panel, lanes 2 and 3). Phospho-transfer was specific for T11, as shown by western blot (**Figure 3B**, middle panel, lanes 2 and 3) and inability to label an H3 1-20 peptide that was already phosphorylated on T11 (**Figure 3B**, lane 5). Interestingly, GST-Mek1 was also unable to phosphorylate a peptide carrying a phosphate on S10 (**Figure 3B**, top panel, lane 4).

The second assay used a semisynthetic epitope system (ALLEN *et al.* 2007) to detect phosphorylation of H3 by Mek1. GST-Mek1 or GST-Mek1-as were incubated with recombinant H3 and the ATPγS analog, 6-Fu-ATPγS. Thiophosphates transferred by Mek1 to substrates were then alkylated to create an epitope that could be detected on western blots with an anti-thiophosphate ester antibody (NIU *et al.* 2009; LO AND HOLLINGSWORTH 2011). Both GST-Mek1 and GST-Mek1-as exhibited autophosphorylation and phosphorylation of H3 (**Figure 3C**, lanes 2 and 5). Moreover, 1-NA-PP1 inhibited both autophosphorylation and H3 phosphorylation by GST-Mek1-as (**Figure 3C**, lane 4), ruling out the possibility of a contaminating kinase phosphorylating H3 T11. We conclude that H3 T11 is a direct substrate of Mek1.

Limitations of a plasmid shuffle system for examining histone mutants

To determine the function of H3 T11 phosphorylation, we constructed strains carrying targeted mutations of T11 alone and in combination with other histone mutations. We initially tested an existing plasmid shuffle system (AHN *et al.* 2005) by porting it to the SK1 strain background. In this approach, also used independently by others (GOVIN *et al.* 2010), the endogenous histone genes were deleted and complemented by wild-type histone genes on a *URA3*-marked *ARS-CEN* plasmid. Histone mutants were introduced on a separate *LEU2 ARS-CEN* plasmid and loss of the *URA3* plasmid was selected for on medium containing 5-FOA.

However, this approach was sub-optimal because of the poor stability of the *ARS-CEN* plasmids in SK1. For example, when liquid cultures of the base histone-deletion strain carrying the *URA3* covering plasmid were grown under conditions selective for the plasmid (i.e., synthetic complete medium lacking uracil), plating on solid medium yielded an efficiency of only 67.2% ± 4.9% (mean ± SD of 5 replicates; colony-forming units per cell plated). Assuming that most cells that failed to form a colony were those that had lost the plasmid because of missegregation during mitosis, it is likely that plasmid copy number per cell is highly variable in the population. Cells with one vs. two copies of an H3/H4-encoding plasmid would likely differ in total histone protein levels and/or have different imbalances with endogenous H2A/H2B. Altered histone gene dosage can cause deleterious effects (MEEKS-WAGNER AND HARTWELL 1986; CLARK-ADAMS *et al.* 1988), so it is possible that cell-to-cell heterogeneity in histone gene copy number might mask or exacerbate the effects of histone point mutations. Furthermore, differences in copy number might have a substantial effect on variation in viability of spores (see below). Finally, although cells in the culture that have lost the histone plasmid would be inviable and therefore presumably would not sporulate, they would contribute to population average measurements in physical assays of recombination.

To circumvent these limitations, we turned to mutagenesis methods that use gene replacement or stable chromosomal integration (Materials and Methods). Stable integration is relatively rapid and obviates concerns about plasmid stability and heterogeneous gene dosage,

but may not fully recapitulate expression from endogenous histone gene loci. The gene replacement strategy provides an even cleaner manipulation of histone genotype, but is more cumbersome because it requires separately mutating two histone gene loci.

Absence of H3 T11 phosphorylation causes little or no overt phenotypes by itself

We replaced both endogenous H3 genes (*HHT1* and *HHT2*) with *hht1-S10A, T11V* and *hht2-S10A, T11V* mutant alleles to eliminate phosphorylation of both S10 and T11. This mutant expressed normal H3 protein levels and neither H3 S10ph nor H3 T11ph could be detected, as expected (**Figure 4A, lanes 3–4**). The mutant displayed normal vegetative growth (**Figure 4B**), similar to a recent report (LI *et al.* 2015). Surprisingly, however, the mutant also displayed normal spore viability (**Table 1**). Meiotic DSBs appeared in normal numbers and locations and disappeared with normal kinetics as assessed by Southern blotting of pulsed-field gels probed for chromosome III (**Figure 4C**), and meiotic progression was not delayed (**Figure 4D**). These results indicate that most if not all meiotic events occur efficiently in the complete absence of both S10ph and T11ph.

To more easily manipulate histone mutants, we used a chromosomal integration strategy to introduce genes for just H3 and H4 as a pair (*HHT2-HHF2*) or all four core histones (*HTA1-HTB1, HHT2-HHF2*) in strains deleted for the endogenous genes for H3-H4 or all four histones. Wild-type or mutant histone genes were integrated on chromosome III at *LEU2*. Strains expressing H3 S10A, T11V, or T11A single mutant proteins or the H3 S10A T11V double mutant were examined in meiotic time courses for H3 S10 and T11 phosphorylation (**Figure 4A**). Importantly, H3 T11 could still be phosphorylated when S10 was mutated to alanine (**Figure 4A, lanes 9–12**); the lower signal in the anti-H3 T11ph western blot could reflect reduced T11 phosphorylation or decreased antibody affinity due to the changed epitope. Similarly, mutation of H3 T11 to alanine or valine did not prevent phosphorylation of S10, as detected with the polyclonal anti-H3 S10ph antibody, although recognition by the monoclonal anti-H3 S10ph antibody was sensitive to these mutations (**Figure 4A, lanes 13–18 and 21–22**).

As with gene replacement, all of these mutants yielded timely meiotic divisions (**Figure 4D**) and spore viabilities indistinguishable from matched wild-type controls (**Table 1**). H3 T11A also supported wild-type interhomolog recombination between *arg4* heteroalleles [23 ± 1.5 Arg^+ recombinants per 1000 viable cells for wild type (SKY3428) vs. 24 ± 0.8 for H3 T11A (SKY3431), mean ± SD for three independent cultures]. Other mutations of H3 T11 yielded similar results: changing T11 to serine or potential phosphomimetic residues (T11D or T11E) again yielded wild-type spore viability (**Table 1**). Mutating H3 T11 also did not reduce spore viability when combined with mutation of H2A S129 [which is also by itself largely dispensable for proper meiosis (SHROFF *et al.* 2004; HARVEY *et al.* 2005)] or with absence of the H3 K4 methyltransferase Set1 [which governs DSB distributions (SOLLIER *et al.* 2004; BORDE *et al.* 2009; ACQUAVIVA *et al.* 2013; SOMMERMEYER *et al.* 2013)] (**Figure 4A, lanes 21-22** and **Table 2**).

Mek1 is required for arrest or delay of meiotic progression when recombination is defective (XU *et al.* 1997; BAILIS AND ROEDER 2000). If H3 T11ph contributes substantially to this Mek1 function, then T11 mutations should alleviate some or all of the meiotic block in *rad51*Δ or *dmc1*Δ mutants. However, in cells lacking Rad51, the *H3 S10A T11V* mutation had negligible effect on either the timing or efficiency of meiotic divisions (**Figure 4E**) and failed to rescue the spore inviability (**Table 2**). This H3 mutation also failed to alleviate the more

stringent arrest in a *dmc1*Δ mutant (**Figure 4E**). Thus, H3 T11ph is dispensable for this checkpoint arrest function of Mek1.

Our findings differ from a prior report of an approximately 35% decrease in spore viability with plasmid-borne *H3 T11A* single or *S10A T11A* double mutants (GOVIN *et al.* 2010). We obtained the published *T11A* plasmid and histone-deleted SK1 host strain (generously provided by J. Govin and S. Berger), verified the *T11A* mutation by sequencing, and carried out the plasmid shuffle. Three independent 5-FOA-resistant clones for each genotype were sporulated and tetrads dissected for wild type and *H3 T11A* side-by-side. The experiment was repeated three times by two investigators. In our hands this *H3 T11A* mutant again yielded spore viability indistinguishable from the control with a wild-type H3 plasmid (**Figure 4F and Table 1**, $p > 0.9$ by linear regression). However, unlike the normal spore viability observed in the stable integrant and gene replacement strains (**Table 1**), viability was consistently lower with plasmid-borne histone genes regardless of H3 genotype (**Figure 4F and Table 1**). A similar defect was reported previously (GOVIN *et al.* 2010). Furthermore, there was substantial heterogeneity in viability from experiment to experiment and between clones within each experiment (**Figure 4F and Table 1**). Within-experiment heterogeneity likely reflects stochastic culture-to-culture variability caused by plasmid instability. Between-experiment variability may reflect differences in sporulation conditions that in turn affect plasmid stability or the sensitivity of these strains to alterations in histone gene expression.

As a counter-example, we also examined a more extreme H3 mutant in which the entire amino-terminal tail was deleted (*H3 ΔN*). The truncated histone was expressed at levels similar to full-length H3 in vegetative cells (**Figure 4A, lanes 23-24**). This mutant displayed vegetative growth defects (**Figure 4B**), delayed and less efficient meiotic divisions (**Figure 4D**), and reduced spore viability (**Table 1**; $p = 0.45$, Fisher's exact test).

H3 T11ph contributes weakly to Mek1 function in the absence of Rad54 T132 phosphorylation

Because *H3 T11* mutations caused no overt defects on their own, we asked whether H3 T11ph might be redundant with Mek1 phosphorylation of Rad54 on T132 (NIU *et al.* 2009). A *rad54-T132A* mutation has little effect by itself, but in a *dmc1Δ* background it allows enough Rad51 activity to partially bypass arrest and produce some viable spores (NIU *et al.* 2009).

In a *rad54-T132A dmc1*Δ background, *H3 T11V* mutation significantly reduced spore viability (**Table 2**; $p = 0.021$, Fisher's exact test), with a decrease in four-spore-viable tetrads and an increase in two- and zero-spore-viable tetrads (**Figure 4G**; $p = 8.1 \times 10^{-5}$, Fisher's exact test). This segregation pattern is diagnostic of increased MI nondisjunction. In this context, *H3 T11V* gave at best only a small increase in overall meiotic division efficiency (**Figure 4E**).

These results suggest that H3 T11 phosphorylation provides a modest contribution to Mek1 function when meiotic recombination defects are encountered. Possible roles of H3 T11ph in these contexts are addressed in the Discussion. However, since the *H3 T11* mutation by itself does not detectably phenocopy a *mek1*Δ mutant, we conclude that H3 T11ph is normally dispensable for Mek1 function.

H3 T11ph enrichment at axis-associated sites and, less strongly, along chromatin loops

H3 T11ph has been used as a cytological marker for Mek1 activity (SUBRAMANIAN *et al.* 2016). Given our results establishing that H3 is a direct phosphorylation target of Mek1, we reasoned that H3 T11ph would also provide a sensitive and specific marker to reveal the

genomic locations of active Mek1 kinase. We therefore assessed H3 T11ph genome-wide by ChIP-seq.

Samples were collected at 3 and 4 hr in meiosis from each of two independent wild-type cultures. To control for specificity of the H3 T11ph ChIP-seq signal, a sample was also collected from a 3.5-hr culture of a *spo11-Y135F* mutant. A set amount of *S. pombe* meiotic cells (4.5 hr in meiosis; 1% of the number of *S. cerevisiae* cells) was added to each *S. cerevisiae* cell sample prior to extract preparation. Mononucleosomes were liberated from formaldehyde-fixed meiotic chromatin by digestion with micrococcal nuclease (MNase), immunoprecipitated with the anti-H3 polyclonal or anti-H3 T11ph monoclonal antibodies, then the DNA was purified and deep sequenced and reads were mapped to the *S. cerevisiae* and *S. pombe* genomes. Each *S. cerevisiae* coverage map was normalized according to *S. pombe* read density for the same antigen from the same culture (**Figure 5A–C** and **Figure S2A,B**). The *S. pombe* spike-in control served two purposes. First, it helped minimize the effects of sample-to-sample variation in lysis, immunoprecipitation, and sequencing library preparation. Second, because the ratio of *S. cerevisiae* to *S. pombe* cells was fixed, the spike-in control provided a scaling factor to compare the relative yield of H3 or H3 T11ph between different *S. cerevisiae* samples (**Figure 5A–C**). Note that this allows comparison between samples for the same antigen, but does not quantify the yields of different antigens relative to one another.

Several lines of evidence establish that these maps reported the distribution of H3 and DSB-dependent H3 T11ph with good specificity. At fine scale, H3 ChIP-seq coverage was low in promoters and showed prominent nucleosome-width peaks in coding sequences (**Figure 5C**), as expected for promoter-associated nucleosome-depleted regions (NDRs) and positioned nucleosomes in gene bodies (JIANG AND PUGH 2009). Replicate samples agreed well, with all five H3 ChIP samples showing highly correlated distributions whether considered genome-wide (**Figure 5D**) or at individual loci (**Figure S2C**). For H3 T11ph, the four wild-type maps correlated well with one another but correlated poorly with either H3 ChIP-seq or H3 T11ph from *spo11-Y135F* (**Figure 5D**), as expected if the ChIP-seq signal was specific for this Mek1-dependent histone modification. Moreover, relative to the *S. pombe* spike-in, *S. cerevisiae* H3 ChIP-seq coverage was similar in wild-type and *spo11-Y135F* samples (**Figure S2C**), but H3 T11ph ChIP-seq coverage was substantially higher in all four wild-type samples than in *spo11-Y135F* (range of 3.4- to 11.6-fold across samples for genome-wide average) (**Figure 5A–C** and **Figure S2D**). The magnitude of the H3 T11ph signal (relative to spike-in) differed by up to ~3.6 fold between the wild-type samples, possibly due to differences in read depth or in culture synchrony or efficiency. Nevertheless, the spatial patterns were highly reproducible (**Figure S2D**), so maps for wild type were averaged for further analysis.

At fine scale, H3 T11ph ChIP coverage showed depletion in NDRs and nucleosomal peaks at similar positions as in the H3 map (**Figure 5C**). This pattern is as expected since presence of a nucleosome (as revealed by bulk H3 localization) is a prerequisite for placement of H3 T11ph by Mek1. However, when maps were examined at larger size scales, H3 T11ph showed broad hills and valleys that were not matched in the H3 ChIP-seq (**Figure 5A,B**), revealing that H3 T11ph tends to be relatively enriched or depleted in domains several kb in width.

A priori, we envisioned two non-exclusive scenarios that might describe H3 T11ph localization: Enrichment at chromosome axes because that is where Mek1 protein is enriched cytologically and Mek1 interacts with axis proteins (BAILIS AND ROEDER 1998; WAN *et al.* 2004;

CARBALLO *et al.* 2008); or enrichment centered on DSB hotspots because Mek1 activation is a response to DSBs and Mek1 regulates DSB repair. We examined each possibility in turn.

Axis-associated sites: To test if H3 T11ph is enriched near axes, we compared its ChIP-seq signal with the genome-wide distribution of axis component Red1 (PANIZZA *et al.* 2011). The sites where ChIP signals for Red1 and other axis proteins are enriched are generally assumed to be the chromatin loop bases that are embedded in the chromosome axis (BLAT *et al.* 2002; PANIZZA *et al.* 2011; SUN *et al.* 2015). These sites often but not always overlap with intergenic regions between convergent transcription units, presumably because transcription can push cohesin and associated axis proteins along chromosomes (LENGRONNE *et al.* 2004; BAUSCH *et al.* 2007; SUN *et al.* 2015).

Across individual chromosomal segments, peaks and valleys in the H3 T11ph signal appeared to correspond well with peaks and valleys of Red1 (**Figure 5A**). Confirming this impression, average H3 T11ph signal formed a broad peak ~4 kb wide when centered on Red1 ChIP-chip peaks, similar in dimensions to the average of Red1 itself and of another axis component, Hop1 (**Figure 5E**). No such enrichment was observed in the *spo11-Y135F* mutant (**Figure 5A,E**). H3 ChIP-seq also showed a weak enrichment centered on Red1 peaks, indicative of a tendency toward higher average nucleosome occupancy, but this was quantitatively modest relative to baseline H3 levels and was similar in wild type and *spo11-Y135F* (**Figure 5E**). Importantly, H3 T11ph signal remained elevated at Red1 peak positions even after correcting for bulk H3 levels (green line in **Figure 5E**). Furthermore, H3 T11ph ChIP-seq correlated well genome-wide with Red1 and Hop1 ChIP signals, whereas H3 ChIP-seq correlated much more weakly (**Figure 5F**). We conclude that H3 T11ph is particularly prevalent where Red1 and Hop1 are enriched, and thus that Mek1 is highly active at axis-associated sites.

Around DSB hotspots: To test if H3 T11ph is enriched near DSB sites, we compared its ChIP-seq signal with DSB maps generated by sequencing of Spo11 oligos (PAN *et al.* 2011; MOHIBULLAH AND KEENEY 2016). When centered on Spo11-oligo hotspots, histone ChIP-seq coverage showed a complex pattern of highly localized enrichment and depletion (**Figure 5G**). The average for total histone H3 showed strong depletion in hotspot centers, flanked by shallow alternating peaks and valleys (gray line in **Figure 5G**). This is the expected pattern from prior studies, reflecting the strong preference for DSBs in *S. cerevisiae* to form in promoter NDRs that are flanked by positioned nucleosomes (OHTA *et al.* 1994; WU AND LICHTEN 1994; PAN *et al.* 2011) (e.g., **Figure 5C**). [For clarity, the plots show averages for the hottest 25% of all hotspots after excluding unusually wide hotspots (> 500 bp); qualitatively similar results were obtained if all hotspots were averaged (data not shown).]

The average H3 T11ph ChIP-seq signal differed from this pattern in informative ways (black line in **Figure 5G**). First, at all positions across the averaging window, H3 T11ph ChIP-seq signal was much higher in wild type than in *spo11-Y135F*, and this difference was greater for stronger hotspots than for weaker ones (**Figure 5G**). Therefore, there is substantial DSB-dependent (thus presumably Mek1-dependent) H3 T11 phosphorylation all across the regions where DSBs usually form, not just at nearby axis sites.

Second, relative to the baseline genomic H3 T11ph signal, there was strong depletion at hotspot centers, indicated by the narrow cleft (~200 bp wide) in the average profile (**Figure 5G**). This cleft corresponded well to the central cleft in the H3 ChIP-seq average, so we infer that this narrow zone of depletion reflects the fact that there are few histones available to be

phosphorylated within the NDRs where hotspots generally occur. There was a peak at hotspot centers when the H3 T11ph levels were normalized to bulk H3 signal (green line in **Figure 5G**). However, there was also a peak when the *spo11-Y135F* map was normalized for bulk H3, thus much or all of this is a DSB-independent signal. This may be a ChIP-seq artifact, or could reflect a low level of Mek1-independent H3 T11ph enriched near promoters (LI *et al.* 2015). [Although gene promoters have lower nucleosome occupancy compared with the rest of the genome, they are not devoid of nucleosomes. For example, some promoters contain positioned, high-occupancy nucleosomes; some contain nucleosomes but only in a fraction of the population; and some contain sub-nucleosomal histone particles (JIANG AND PUGH 2009; FLOER *et al.* 2010; WEINER *et al.* 2010).]

Third, there was a broader zone of lower H3 T11ph signal flanking the central NDR and extending ~2 kb on either side (**Figure 5G**). This zone extended into areas where bulk H3 levels were high, so the difference map (normalizing H3 T11ph to H3) revealed depletion for H3 T11ph relative to immediate surroundings (green line in **Figure 5G**). Nonetheless, the H3 T11ph signal across this region was substantially higher in wild type than in *spo11-Y135F*. Much of this depleted zone corresponded to the same areas covered by exonucleolytic resection tracts as measured by S1-seq (MIMITOU *et al.* 2017) (blue line in **Figure 5G**). This suggests that some or all of this depletion reflects disruption of chromatin — and thus of ChIP-detectable H3 T11ph signal — accompanying DSB resection. Interestingly, the H3 T11ph depletion zone correlated precisely with the dimensions of a zone of relative depletion for Red1 (**Figure 5G**).

Taken together, these findings suggest that the distribution of DSB-provoked H3 T11 phosphorylation is governed largely by the distribution of Red1 and other proteins that are directly involved in Mek1 activation. Further implications of these patterns are addressed in the Discussion.

H3 T11ph correlates with DSB frequency across large sub-chromosomal domains

We next examined larger scale variation in H3 T11ph ChIP signal across chromosomes. H3 T11ph ChIP signals were binned in non-overlapping windows of varying sizes from 0.5 to 40 kb, then compared (Pearson's *r*) to Spo11-oligo densities or ChIP signals for Red1, Hop1, or Rec8 in the same bins (**Figure 6**). Comparisons using the ratio of H3 T11ph to H3 show which correlations are specific for the histone modification ChIP per se (green points in **Figure 6**) as opposed to underlying (background) enrichment or depletion in the bulk chromatin map (total H3; gray points). Comparisons using the ratio of wild type to *spo11-Y135F* for H3 T11ph show which correlations are specific for DSB-dependent (and thus Mek1-dependent) signal (blue points in **Figure 6**).

For small windows (< 2 kb), both H3 and H3 T11ph were anticorrelated with Spo11-oligo density (**Figure 6A**). This pattern is driven by strong preference for DSBs to form in NDRs, and the attendant depletion of histone signal around hotspots (**Figure 5G**). In contrast, with large windows the H3 T11ph signal instead had a significant positive correlation with Spo11-oligo density, with Pearson's *r* values high over a range of ~25–40 kb (**Figure 6A**). This correlation was also high when the wild-type H3 T11ph ChIP data were normalized to coverage in *spo11-Y135F*, but no such correlation was seen for total histone H3, thus this pattern is specific for DSB-dependent H3 T11 phosphorylation. We infer that subchromosomal domains tens of kb wide that experience more DSBs also incur more Mek1 activity on average. This finding fits with the expectation that H3 T11ph is a faithful molecular reporter of DSB-provoked Mek1 kinase activity.

In contrast to the wide variation in correlation behavior depending on window size when H3 T11ph was compared Spo11-oligo density, comparisons with either Red1 or Hop1 ChIP showed strong positive correlations over all window sizes tested (**Figure 6B,C**). For Rec8, H3 T11ph showed a modest positive correlation for short windows but little or no correlation with larger windows (**Figure 6D**). These patterns can be understood as the combination of two spatial correlations with different length dependencies. At short distances (<10 kb), Mek1 activity is particularly enriched at preferred binding sites for Red1, Hop1, and Rec8 (i.e., axis sites; **Figure 5E**). At longer distances (tens of kb), the domains that are relatively DSB-rich (and thus have more Mek1 activity) are also enriched for Red1 and Hop1 but not for Rec8 (BLAT *et al.* 2002; PAN *et al.* 2011; PANIZZA *et al.* 2011).

DISCUSSION

This study and others (GOVIN *et al.* 2010; SUBRAMANIAN *et al.* 2016) establish that H3 T11 phosphorylation is highly induced during meiosis in *S. cerevisiae*. Our findings additionally demonstrate that H3 T11ph is a direct product of DSB-induced activation of Mek1. Mek1 is conserved in *S. pombe* (PEREZ-HIDALGO *et al.* 2003), so it seems likely that this kinase is also responsible for the H3 T11ph we observed in fission yeast.

Mek1 appears specifically in fungal taxa, but the larger Rad53 kinase family is ubiquitous in eukaryotes (SUBRAMANIAN AND HOCHWAGEN 2014). Another member of this family, CHK1, was reported to be required for H3 T11ph in mouse fibroblasts (SHIMADA *et al.* 2008). In this case, however, DNA damage caused a decrease in H3 T11ph levels. It remains unknown if CHK1 directly phosphorylates H3 T11 or if H3 T11ph occurs in response to DSBs in mammalian meiosis. H3 T11ph has been reported during meiosis in sciarid flies (ESCRIBA *et al.* 2011), suggesting evolutionary conservation beyond yeasts.

H3 T11 can also be directly phosphorylated by pyruvate kinase M2 in *S. cerevisiae* and mammalian cells, possibly to coordinate chromatin structure and gene expression with the cell's nutritional status (YANG *et al.* 2012; LI *et al.* 2015). In cultured human cells, H3 T11ph is also formed by protein-kinase-C-related kinase 1 near promoters of androgen receptor-modulated genes (METZGER *et al.* 2008), and by death-associated protein (DAP)-like kinase during mitosis, particularly near centromeres (PREUSS *et al.* 2003). Our results establish that meiotic induction of H3 T11ph in yeasts is fundamentally distinct from these other modes of H3 T11 phosphorylation in terms of provenance and genomic distribution.

Possible functions of H3 T11ph in meiosis

Under the conditions in this study, histone mutations that eliminated H3 T11 phosphorylation caused no discernible meiotic defects by themselves. This was true with multiple independent mutagenesis strategies and numerous mutant constructs encoding different amino acid substitutions alone or in combination with mutation of H3 S10. We conclude that H3 T11ph is dispensable for meiosis under our standard conditions.

Why our results differed from a previous report (GOVIN *et al.* 2010) remains unknown. One possibility is that the highly variable spore viability in the plasmid shuffle system fortuitously gave the incorrect appearance of a meiotic defect in the earlier study. The reported decrease in spore viability [from ~80% in the control to ~50% with *H3 T11V* (GOVIN *et al.* 2010)] was of comparable magnitude to the intrinsic experimental variability we observed with plasmid-borne histone cassettes. Alternatively, studies in the two laboratories may have had undocumented differences in sporulation conditions to which *H3 T11* mutants are specifically sensitive.

Despite H3 T11ph being dispensable in unperturbed meiosis, we did observe that blocking phosphorylation of H3 T11 modestly exacerbated the phenotype of a *dmc1*Δ *rad54-T132A* mutant. One interpretation is that H3 T11ph helps Mek1 maintain residual interhomolog bias when Rad51 is the sole source of strand exchange activity. In this model, increased MI nondisjunction is caused by more of the residual DSB repair being between sister chromatids, and less between homologs. This interpretation is motivated by the increased intersister recombination observed in a *rad54-T132A* mutant when Mek1 activity is inhibited, and by the ability of the *rad54-T132A* mutation to rescue some spore viability in a *dmc1*Δ background but not in *dmc1*Δ *mek1*Δ (NIU *et al.* 2009). These findings indicated that other Mek1 targets contribute to interhomolog recombination by Rad51 when Dmc1 is missing and Rad54 cannot be

phosphorylated. The recent discovery that Mek1 phosphorylates Hed1 and histone H2B make these strong candidates for additional redundancy (CALLENDER *et al.* 2016; SUHANDYNATA *et al.* 2016) (N.M.H., unpublished data).

If H3 T11ph does promote Mek1 function, albeit in a minor way, what might its role be? One possibility is that it is an effector of Mek1 signaling. This could be via recruitment to chromatin of proteins with phosphothreonine binding motifs such as the FHA domain, which is present in numerous proteins in *S. cerevisiae* including the recombination protein Xrs2 (MAHAJAN *et al.* 2008; MATSUZAKI *et al.* 2008). Or, H3 T11ph might impinge on nucleosome stability, higher-order chromatin organization, or ability to install, remove, or read other histone modifications. We observed potential crosstalk between histone modifications in that H3 S10ph blocked the ability of Mek1 to phosphorylate T11 on the same peptide. Crosstalk of H3 T11ph with other H3 modifications has been documented in vegetatively growing yeast [H3 K4 methylation (LI *et al.* 2015)] and in human cells [H3 K9 acetylation (YANG *et al.* 2012) and demethylation (METZGER *et al.* 2008)]. A second, non-exclusive possibility is that H3 T11ph might maintain or amplify Mek1 activity via positive feedback. For example, the FHA domain of Mek1 might bind directly to H3 T11ph in a manner that stabilizes or increases the amount of active Mek1. Both general types of role — downstream effector or feedback amplifier — are compatible with the observed genetic interaction of *H3 T11* mutation with *dmc1*Δ *rad54-T132A*.

Spatial organization of Mek1 activity

Although H3 T11 can apparently be phosphorylated by other kinases, the magnitude of the DSB- and Mek1-dependent signal combined with its rapid disappearance when Mek1 is shut off made H3 T11ph an excellent candidate for a molecular marker of ongoing Mek1 activity. Our experiments establish proof of principle for this use in genomic experiments, and also validate that cytological staining for H3 T11ph provides a direct readout of Mek1 activity (SUBRAMANIAN *et al.* 2016).

The most prominent sites of H3 T11ph, and thus of Mek1 activity, were coincident with peaks of Red1 and Hop1, i.e., presumed axis attachment sites. This pattern is not surprising given that Mek1 protein appears to be enriched on axes as assessed by immunocytology (BAILIS AND ROEDER 1998). However, immunolocalization does not reveal kinase activity per se, and cannot evaluate the degree to which activity might spread in *cis*. Interestingly, the H3 T11ph enrichment was highly similar to that of Red1 and Hop1 around axis sites. This more localized pattern contrasts with the spread of γ-H2A over tens of kb around DSBs in yeast (SHROFF *et al.* 2004). This pattern could be because Mek1 protein is constrained, i.e., it rarely diffuses away from the sites where it has been activated. Alternatively, Mek1 may be rapidly inactivated if it diffuses away and/or the phosphates that Mek1 places outside the immediate vicinity of Mek1 activation sites might be more rapidly removed by phosphatases.

However, substantial levels of DSB-dependent H3 T11ph were also observed across areas in between Red1 peaks, i.e., along the lengths of presumptive chromatin loops. This signal correlated with Spo11-oligo frequency (i.e., local DSB density) but was lower immediately around DSB hotspots, coincident with regions also depleted of Red1. For any protein, ChIP enrichment at given sites does not imply that these are the only sites of binding. In yeast meiosis, axis proteins Red1 and Hop1 are also detected above background across regions in between the sites of their principal enrichment. ChIP provides a population average measurement, so possible interpretations are that Red1 et al. are bound to different sites in different cells, but only at discrete sites in any one cell (an "axis sites only" model); or that there is lower level binding to

chromatin loops in addition to higher level binding at axis sites (an "axis plus dispersed binding" model).

The DSB-dependent H3 T11ph ChIP signal tracked closely with Red1 and Hop1 ChIP at all size scales, both quantitatively and spatially. Thus, our findings are consistent with Mek1 activity manifesting anywhere Red1 and Hop1 are present to support it once DSBs have formed. It is interesting to note that the sites of highest Mek1 activity (i.e., Red1/Hop1 peaks) are spatially distinct from the DSB sites where Mek1 exerts its known biological function — Hop1/Red1-dependent control of recombination outcome. However, even though H3 T11ph ChIP signal was less abundant immediately adjacent to DSB sites than elsewhere, the DSB-proximal signal was substantially higher in wild type than in the *spo11-Y135F* control. Thus, we infer that active Mek1 kinase has access to chromatin and chromatin-associated proteins immediately surrounding DSBs.

A puzzle about the signal immediately adjacent to hotspots is that DSBs are exonucleolytically resected for ~800 nucleotides on average on both sides of the break (ZAKHARYEVICH *et al.* 2010; MIMITOU *et al.* 2017), but ssDNA should not be revealed in our ChIP-seq data even if it were still bound by histones because the sequencing library preparation protocol is not expected to support recovery of ssDNA. What then is the source of H3 T11-phosphorylated nucleosomes immediately around hotspots? Likely candidates are the sister of the broken chromatid, one or both intact chromatids of the homologous chromosome with which recombination is occurring, and/or recombination intermediates (D-loops and double Holliday junctions) assuming that these are chromatinized. Because Mek1 controls homolog bias, we speculate that some or all of the H3 T11ph signal around hotspots is from Mek1 action on the sisters of broken chromatids. In support of this idea, we further note that the areas expected to be covered by DSB resection are also areas where the H3 T11ph signal is locally depleted. This pattern is thus in line spatially with what would be predicted if both sister chromatids are being exposed to Mek1 activity.

Our data are consistent with spreading of DSB-provoked Mek1 activity in *cis* along chromatin and concentrating wherever Red1 and Hop1 are also concentrated. The findings neither refute nor support the TLAC model (see Introduction), but are consistent with this model provided that DSB-dependent activation of Mek1 at axis sites can be accompanied by spreading of Mek1 activity to surrounding chromatin as well. Current versions of the TLAC model favor the idea that tethering occurs before DSB formation because some partners of Spo11 are enriched at axis sites rather than at hotspots but can be connected to hotspots physically via interactions with a reader (Spp1) of the H3 K4 methylation that is prominent around promoters (PANIZZA *et al.* 2011; ACQUAVIVA *et al.* 2013; SOMMERMEYER *et al.* 2013). Such loop-axis interactions prior to DSB formation could provide a means to rapidly and specifically activate Mek1 at a nearby axis site in response to a DSB at a hotspot within a tethered loop.

Immunostaining experiments demonstrated H3 T11ph foci of variable size and intensity on chromatin where synaptonemal complex had not yet formed, although spatial disposition of this phosphorylation relative to axes has not been reported (SUBRAMANIAN *et al.* 2016). Combining these cytological findings with our genomic data suggests that each DSB provokes a relatively large zone of Mek1 activation that encompasses the broken chromatin loop, its sister chromatid, and the adjacent loop bases(s). We speculate that this zone may extend across one or more loops, perhaps dependent in part on availability of sufficient amounts of Red1 and Hop1 to support Mek1 activity.

In summary, the detection of H3 T11ph is useful as an indicator of meiotic DSB formation, an indicator of Mek1 activation level, and a marker of the spatial organization of chromatin that Mek1 acts upon. H3 T11ph ChIP will be a powerful tool for dissecting not only the function of Mek1 but also the higher order structural organization of recombining chromosomes.

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FIGURE LEGENDS

Figure 1. H3 T11 phosphorylation in *S. cerevisiae* meiosis.

(A) Western blots of whole-cell extracts from asynchronous cycling vegetative (Cyc) and synchronized meiotic culture time points in wild-type and mutant strains. In panels i–iv, the antibodies used were anti-H3 T11ph polyclonal (Active Motif 39151), anti-H3 S10ph monoclonal (EMD Millipore 05-817), anti-H2A S129ph/γ-H2A (Abcam 15083), and anti-H3 (Abcam 1791). For panels v and vi, anti-H3 T11ph monoclonal (EMD Millipore 05-789) and anti-H3 S10ph polyclonal (EMD Millipore 06-560) were used; other antibodies were the same. Interstitial lanes were removed from the blot images in panel vi to match time points in other panels. Filled and open arrowheads indicate 20 and 15 kDa molecular weight markers, respectively. **(B)** Western blot comparison of anti-H3 T11ph monoclonal (mAb; EMD Millipore 05-789) and polyclonal (pAb; Active Motif 39151) antibodies. **(C)** Meiotic progression assessed by DAPI staining. Cells with ≥2 DAPI-staining bodies were scored as having progressed past the first meiotic division; n ≥ 100 cells per time point. The *rad50S* culture was not quantified past 6 hr because of nuclear fragmentation. **(D)** The first twenty amino acids in histone H3 and modifications known to occur in *S. cerevisiae* or *S. pombe*: ac, acetylation; me, methylation; ph, phosphorylation. **(E)** Meiosis-specificity of DNA damage-induced H3 T11ph. Asynchronous vegetative cultures of wild type were treated with genotoxins that induce DSBs, then whole-cell extracts were prepared and analyzed by western blotting for H3 T11ph. Cultures in the left panel were untreated (Mock) or treated with X-rays (400 Gy) or camptothecin (20 μM) at room temperature. An interstitial lane was deleted from the blot image for this panel. Cultures in the right panel were untreated or treated with X-rays (400 Gy) on ice. Premeiotic (0 hr) and meiotic (4 hr) cultures were included as controls. The anti-H3 T11ph monoclonal (EMD Millipore 05-789) was used. Arrowheads are as defined in panel A.

Figure 2. H3 T11 phosphorylation in *S. pombe* meiosis.

Western blots of whole-cell extracts from haploid *pat1-114* strains undergoing synchronized meiosis. Antibodies used were the same as in **Figure 1Av**. Filled and open arrowheads indicate 20 and 15 kDa molecular weight markers, respectively. The altered electrophoretic mobility of histones at later time points in some cultures was probably caused by varying levels of contaminating DNA in the extracts rather than differential post-translational modifications.

Figure 3. H3 T11 is a direct target of Mek1 kinase.

(A) Persistence of H3 T11ph requires maintenance of Mek1 kinase activity. A meiotic culture of a *mek1-as*, *dmc1Δ* (strain SKY3095) was split 4 hr after transfer to sporulation medium. One part was left to continue in meiosis untreated, the other part was treated with 1 μM 1-NA-PP1. Whole-cell extracts were prepared at the indicated times and assayed for H3 T11ph by western blotting (mAb; EMD Millipore 05-789). Filled and open arrowheads indicate 20 and 15 kDa molecular weight markers, respectively. Numbers indicate hours after transfer to sporulation medium. **(B)** Mek1 kinase assay using radioactive ATP. Affinity-purified GST-Mek1 (250 ng) was incubated in the presence of [γ-^{32}P]ATP either alone or with 2 μg recombinant H3 or 5 μg of unphosphorylated or phosphorylated synthetic H3 1-20 peptides as substrates. Reactions were separated by SDS-PAGE and visualized by autoradiography (top), anti-H3 T11ph western blot (middle; polyclonal Active Motif 39151), and Coomassie staining. **(C)** Mek1 kinase assay by semisynthetic epitope labeling. Kinase reactions were carried out with affinity-purified GST-Mek1 (2 μg) or GST-Mek1-as (0.76 μg) in the presence of ATPγS or 6-Fu-ATPγS with 2 μg

recombinant H3. After incubation 30 min at 30°C, PNBM (*p*-nitrobenzyl mesylate) was added to alkylate the thiophosphorylated target sites. Reactions were then separated by SDS-PAGE and analyzed by western blotting with anti-thiophosphate ester monoclonal antibody (top panel; Epitomics 2686-1) or anti-H3 T11ph monoclonal antibody (EMD Millipore 05-789). Interstitial lanes were removed from images in panels B and C as indicated by the white lines.

Figure 4. Characterization of histone mutant strains.
(A) Composite of western blots of whole-cell extracts from synchronous meiotic cultures or asynchronous cycling vegetative cultures ("C") carrying the indicated histone mutations. Antibodies used were: anti-H3 T11ph polyclonal (Active Motif 39151) or anti-H3 T11ph monoclonal (EMD Millipore 05-789); anti-H3 S10ph monoclonal (EMD Millipore 05-817); anti-H3 S10ph polyclonal (EMD Millipore 06-560); anti-γ-H2A (Abcam 15083); and anti-H3 (Abcam 1791). Filled and open arrowheads indicate 20 and 15 kDa molecular weight markers, respectively. "n.d." indicates not determined; "*hht2-ΔN*" encodes H3 lacking its N-terminal 30 amino acids. **(B)** Vegetative growth of H3 mutant strains. Cells from overnight cultures were spotted onto YPD plates using a manifold pin replicator and represent 1:5 serial dilutions starting with ~2.5×10^6 cells/ml. **(C)** Analysis of meiotic DSB formation. High-molecular-weight DNA isolated in agarose plugs was separated by pulsed-field gel electrophoresis followed by Southern blotting and indirect end-labeling with a probe directed against *CHA1* on the left arm of chromosome III. The lower panel shows quantification of the DSB signal as percent of lane total after background subtraction. (**D,E**) Meiotic progression of representative histone mutant strains. Cells were fixed and stained with DAPI and the fraction of cells with ≥ 2 nuclei was counted (n ≥ 100 cells per time point). For panel E, strains used were *rad51Δ* (SKY3183); *rad51Δ, H3 S10A, T11V* (SKY3186); *dmc1Δ, rad54 T132A* (SKY3802); *dmc1Δ, rad54 T132A, H3 T11V* (SKY3659); *dmc1Δ* (SKY3078) and; *dmc1Δ H3 S10A, T11V* (SKY3091). **(F)** Spore viabilities in plasmid shuffle strains expressing wild-type H3 or H3 T11A. Three independent clones isolated for each genotype were sporulated and tetrads were dissected in three separate experiments. Each point represents the value from a single isolate (n = 30–32 tetrads per data point). See **Table 1** for summary and text for statistical test. Strains used were: H3 wild type (SKY3438-3440) and *H3 T11A* (SKY3441-3443). **(G)** Evidence that the *H3 T11V* mutation increases MI nondisjunction in a *rad54-T132A dmc1*Δ background. The distribution of viable spores in tetrads is shown for the indicated strains. An increase in 2- and 0-spore-viable tetrads (rather than 3- or 1-spore-viable) is diagnostic of an increased frequency of MI nondisjunction. Strains were *dmc1Δ, rad54 T132A* (SKY3802) and *dmc1Δ, rad54 T132A, H3 T11V* (SKY3659).

Figure 5. Spatial disposition of H3 T11ph along meiotic chromosomes.
(**A–C**) Anti-H3 and anti-H3 T11ph ChIP-seq coverage across representative genomic regions. Coverage data for each chromosome were normalized ("norm") relative to an *S. pombe* spike-in control. The four wild-type samples were averaged; the single *spo11-Y135F* sample (denoted "*spo11yf*") is presented separately. For a given antigen (H3 or H3 T11ph), use of the internal *S. pombe* control allows direct quantitative comparison of relative yield in different samples, revealing in particular the DSB-dependent H3 T11ph signal via comparison of wild type with *spo11-Y135F*. The Spo11-oligo map (RPM, reads per million) (MOHIBULLAH AND KEENEY 2016) and decile-normalized anti-Red1 ChIP-chip data (PANIZZA *et al*. 2011) are shown for comparison. The H3 and H3 T11ph data shown in A and B and all Red1 data were smoothed with 500 bp Parzen (triangular) sliding window. Color coding is retained in the other panels in

this figure. (**D**) Reproducibility of histone ChIP-seq coverage maps. ChIP-seq coverage was averaged in 500 bp windows and compared between datasets. The heatmap is shaded according to the Pearson's *r* value for each pairwise comparison. (**E**) H3 T11ph enrichment around presumed axis-attachment sites. H3 (upper graph) and H3 T11ph ChIP-seq coverage (middle graph) and smoothed (500-bp Parzen window) Red1 and Hop1 ChIP-chip data (lower graph, PANIZZA *et al*. 2011) were averaged around 1869 Red1 ChIP peaks. The green lines in the lower graph show the ratio of H3 T11ph to H3 ChIP-seq. (**F**) H3 T11ph correlates well with Red1 and Hop1 ChIP signal genome wide. Each point compares the H3 or H3 T11ph ChIP-seq coverage in wild type with Red1 or Hop1 ChIP-chip signal averaged across non-overlapping 5-kb bins. Correlation coefficients (Pearson's *r*) are indicated in each plot. (**G**) H3 T11ph around DSB hotspots. ChIP-seq and Spo11-oligo data were averaged around Spo11-oligo hotspots and plotted as in panel E (MOHIBULLAH AND KEENEY 2016); for clarity, hotspots more than 500 bp wide were excluded, and only the hottest 25% of hotspots are shown (N=872 hotspots). Note that vertical and horizontal scales for ChIP-seq data are the same in panels E and G to facilitate direct comparison. The blue line in the third panel shows the extent of dsDNA depletion predicted from S1-seq mapping of DSB resection tracts around the same hotspots (MIMITOU *et al*. 2017). The lowest panel shows the average Spo11-oligo profile.

Figure 6. Scale-dependent correlations of H3 T11ph with chromosomal features.
Anti-H3 (gray points) and anti-H3 T11ph (black points) ChIP-seq coverage was binned in non-overlapping windows of varying sizes and compared (Pearson's *r*) to Spo11-oligo density (A) or ChIP-chip signal for Red1 (B), Hop1 (C), or Rec8 (D) averaged across the same windows (ChIP-chip data from, PANIZZA *et al*. 2011). Green points show correlations using the ratio of H3 T11ph to H3 in wild type; blue points show correlations for the ratio of wild type to *spo11-Y135F* H3 T11ph signal.

SUPPLEMENTAL FIGURE LEGENDS

Supplemental Figure S1. Specificity of anti-H3 T11ph and anti-H3 S10ph antibodies. (**A**) Histone peptide array western blots showing the specificity of (**i**) anti-H3 T11ph mAb or (**ii**) anti-H3 S10ph pAb and their tolerance of neighboring modifications. Blots are of duplicate 384-peptide arrays (MODified Histone Peptide Array, Active Motif 13001) of immobilized synthetic histone H2A, H2B, H3 and H4 unmodified peptides or peptides containing from one to four modified residues including many possible combinations of histone modifications that are found in higher eukaryotes, of which only a small number are known to be present in yeast. Positions A1–L11 contain H3 peptides, L12–O11 contain H4 peptides, O12–P3 contain H2A peptides and P4–P19 contain H2B peptides. Peptides that were highly reactive with either antibody are listed below the blot image; the entire table of peptides is listed in **Supplemental Table S2.** (**B**) Immunodetection of histone H3 amino-terminal peptides (residues 1–20) or recombinant histone proteins spotted onto PVDF membranes demonstrating the specificity of antibodies to phospho-H3 T11 and phospho-H3 S10. Spots were 10-fold serial dilutions of peptides or recombinant histones starting with 167 ng in the left-most column. Recombinant histone proteins produced in *E. coli* were from the following species: H2A, H2B, H3 from *S. cerevisiae*; H3.3 from *H. sapiens*; and H4 from *X. laevis*. Antibodies were: anti-H3 pAb (Abcam 1791), which is specific to the carboxy-terminal 35 amino acids of histone H3; anti-H3 T11ph polyclonal (Active Motif 39151); anti-H3 S10ph monoclonal (EMD Millipore 05-817); and anti-H3 S10ph polyclonal (EMD Millipore 06-560).

Notes:

(**Panel Ai**) The diphosphorylated H3 1–19 S10ph T11ph peptide at position D5 was not detected by the anti-H3 T11ph mAb, whereas all phospho-T11 containing peptides (except those that also contained phospho-S10) were detected (peptides containing phospho-T11 along with methyl-K4 were not included in the array). We conclude that this mAb detects only the monophosphorylated peptide, but that it is tolerant of other modifications of the H3 N-terminal tail.

(**Panel Aii**) The diphosphorylated (S10ph T11ph) peptide at position D5 was not detected by the anti-H3 S10ph pAb, whereas all phospho-S10 containing peptides (except those that also contained phospho-T11) were detected (peptides containing phospho-S10 along with methyl-K4 were not included in the array). We conclude that this pAb detects only the monophosphorylated peptide, but that it is relatively tolerant of other modifications of the H3 N-terminal tail. This pAb showed detectable cross-reaction to other modifications as well. Peptides that scored as weakly reactive with anti-H3 S10ph pAb were: J6, H3 1–19 S10ph K14ac; J11, H3 7–26 K18ac; J13, H3 7–26 K14ac R17me2s; J15, H3 7–26 R17me2s K18ac; J19, H3 7–26 K14ac R17me2a K18ac; K4, H3 16–35 S28ph; L7, H3 26–45 unmodified; L8, H3 26–45 K36me1; L9, H3 26–45 K36me2; L11, H3 26–45 K36ac; M18, H4 11–30 unmodified; M19, H4 11–30 K12ac; M20, H4 11–30 K16ac; M21, H4 11–30 R17me2s; M22, H4 11–30 R17me2a; N5, H4 11–30 R24me2a; N6, H4 11–30 R24me2s; N7, H4 11–30 K12ac K16ac; N8, H4 11–30 K16ac R17me2s; N9, H4 11–30 K16ac R17me2a.

(**Panel B**) Anti-H3 T11ph pAb was capable of detecting phospho-T11 even with nearby methylation at lysine 9, a modification that occurs in *S. pombe* and metazoans, but not in *S.*

cerevisiae. Both anti-H3 S10ph antibodies also reacted slightly with full-length recombinant H3 and H3.3.

Supplemental Figure S2. Anti-H3 and anti-H3 T11ph ChIP-seq.
(A) *S. pombe* read coverage. The read coverage from a representative sample (H3 ChIP-seq from wild type culture 1 at 3 hr) is shown. The zones of exceptionally high coverage at the ends of chromosome III are the ribosomal DNA repeats. Note that the read density is too sparse to allow investigation of H3 or H3 T11ph patterns across the *S. pombe* genome. **(B)** Reproducibility of read depth for *S. pombe* chromosomes I and II. Stacked bars show the contribution of reads from chromosomes I and II relative to the total for these two chromosomes. Lower case letters are sample identifiers as indicated in **Figure 5D**. Chromosome III had more variable read coverage (data not shown), possibly because of greater sampling error attributable to presence of the large rDNA arrays. Because relative read depths were stable for chromosomes I and II, these were combined to calculate the normalization factors for the *S. cerevisiae* maps. **(C,D)** Reproducibility of spatial patterns for anti-H3 and anti-H3 T11ph ChIP-seq. The map for each wild-type sample (solid lines) is shown separately, superimposed on the *spo11-Y135F* map (dashed lines) for comparison. The maps depict the same region shown in **Figure 5C**. Note that the H3 ChIP-seq maps were highly reproducible across all four wild-type samples as well as the *spo11-Y135F* sample. For H3 T11ph, although the magnitude of the signal varied between wild-type samples, their spatial patterns were highly similar and all gave substantially higher coverage than the *spo11-Y135F* sample.

Table 1. Absence of H3 T11ph does not compromise spore viability.

Mutation method [a]	H3 genotype [b]	Strain	Spore viability (no. of tetrads)
Replacement	Wild type	2701	97% (44)
	S10A T11V	2705	97% (44)
H3-H4 integration	Wild type	3311	96% (22)
	S10A	3333	97% (64)
	T11V	3342	95% (64)
	S10A T11V	3334	93% (64)
	T11A	3312	95% (86)
	T11S	3313	99% (22)
	T11D	3332	97% (64)
	T11E	3303	97% (64)
Four-core integration	Wild type	3330	94% (22)
	T11V	3264	97% (64)
	Δ*N*	2388	81% (44)
Plasmid shuffle, Expt. 1 [c]	Wild type	3438–3440	90% (90)
	T11A	3441–3443	86% (90)
Plasmid shuffle, Expt. 2 [c]	Wild type	3438–3440	75% (92)
	T11A	3441–3443	77% (94)
Plasmid shuffle, Expt. 3 [c]	Wild type	3438–3440	74% (96)
	T11A	3441–3443	76% (96)

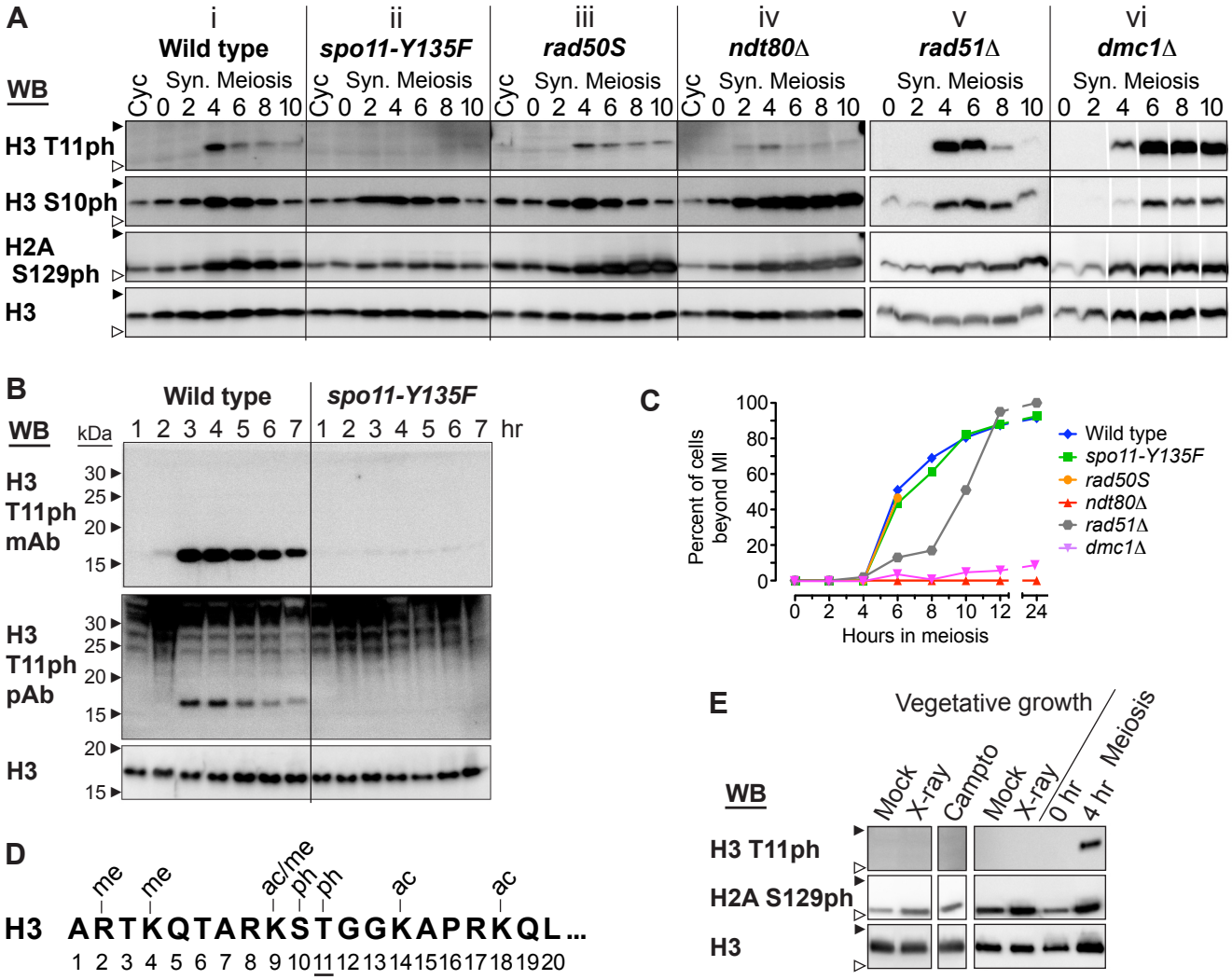
[a] Replacement: Both *HHT1* and *HHT2* were replaced with wild-type or mutant copies at their endogenous locations. Integration: Stable integration at *leu2::hisG* of a cassette carrying either the H3 and H4 gene pair *HHT2-HHF2*, or all four core histone genes *HTA1-HTB1* and *HHT2-HHF2*. Endogenous loci (encoding H3 and H4 or all four core histones, respectively) were deleted. Plasmid shuffle: Replacement of a *URA3* plasmid carrying wild-type *HHT2-HHF2* with a *LEU2* plasmid carrying either wild-type or mutant versions. The endogenous H3 and H4 loci were deleted.
[b] Genotypes are homozygous unless plasmid-based. See **Supplemental Table S1** for complete genotypes.
[c] Three independent 5-FOA^R colonies were isolated for each shuffle plasmid and were dissected separately. The dissections were performed on three separate occasions by two different investigators; all six strains were dissected in parallel in each experiment. A breakdown of results by strain and experiment is provided in **Figure 4F**. Neither histone H3 genotype nor clone identity was a significant predictor of altered spore viability ($p \geq 0.9$, linear regression).

Table 2. Combining *H3 T11* mutations with other mutations.

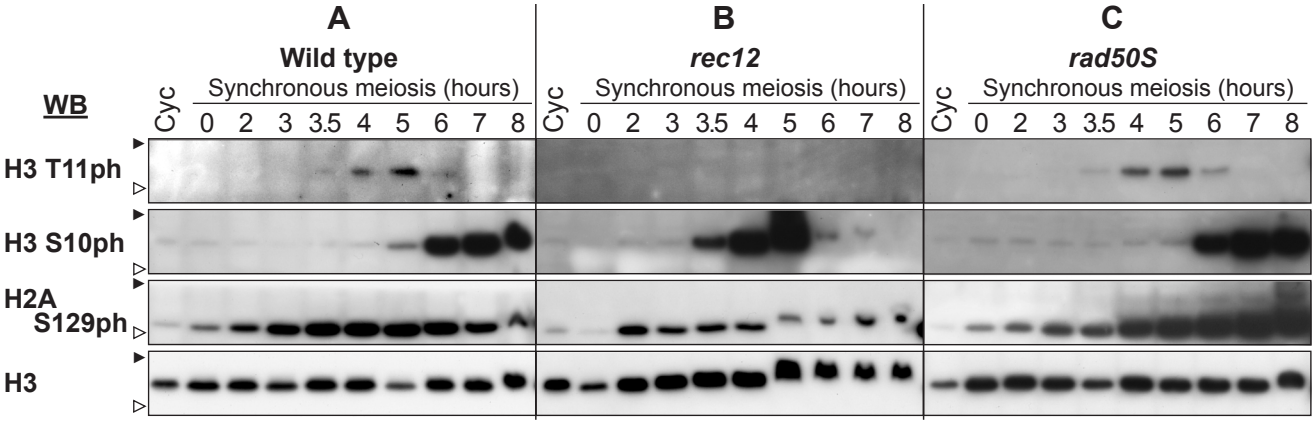
Additional mutation(s)[a]	H3 genotype[a]	Strain	Spore viability (no. of tetrads)
H2A S129A	wild type	3265	97% (64)
	T11V	3331	95% (64)
*set1*Δ	wild type	4415	97% (42)
	S10A T11V	3329	97% (64)
*rad51*Δ	wild type	3183	0% (44)
	S10A T11V	3186	0% (44)
*rad54-T132A dmc1*Δ	wild type	3802	67% (65)
	T11V	3659	49% (86)

[a] Genotypes are homozygous. See **Supplemental Table S1** for complete genotypes.

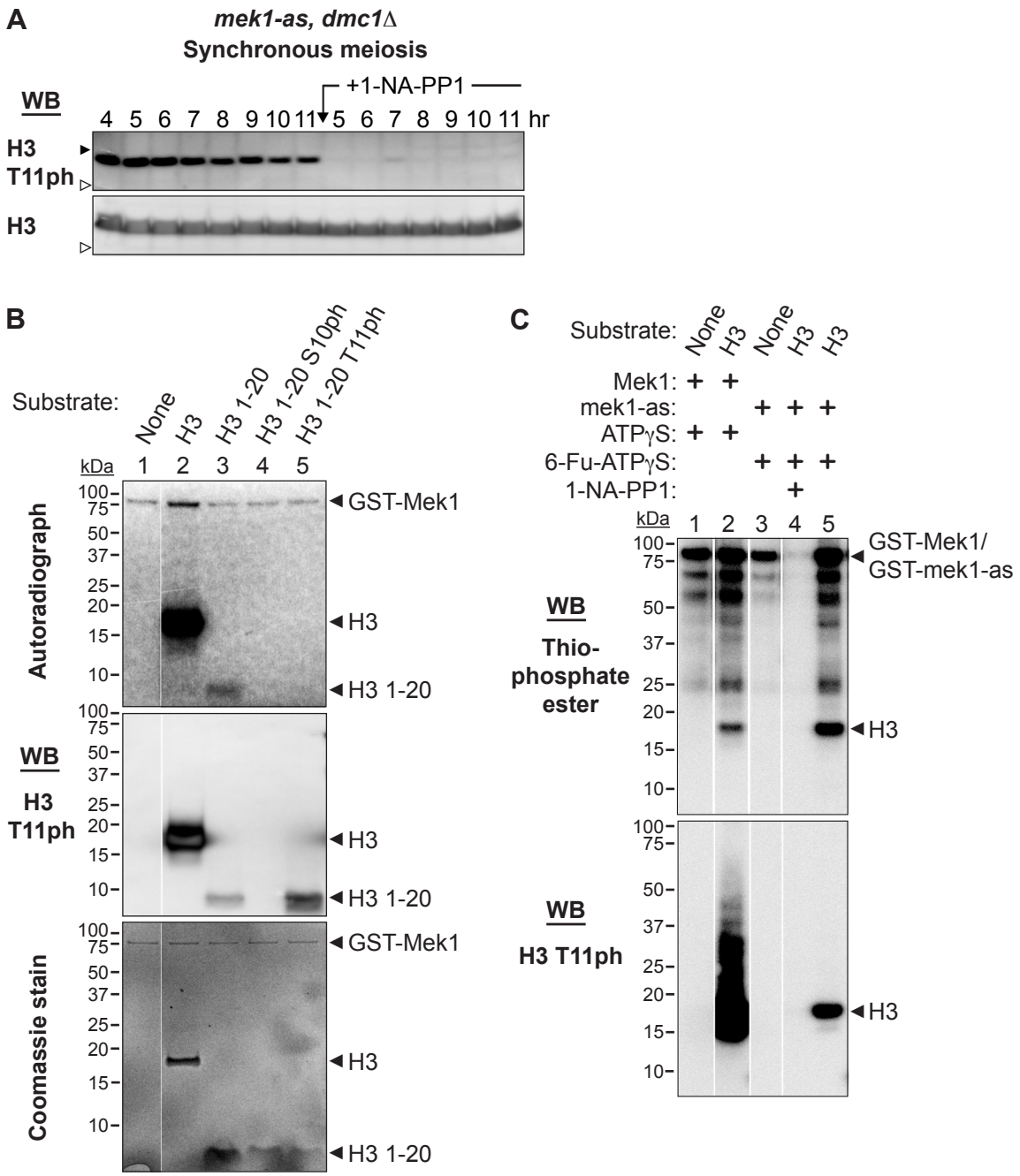
Kniewel et al., Figure 1



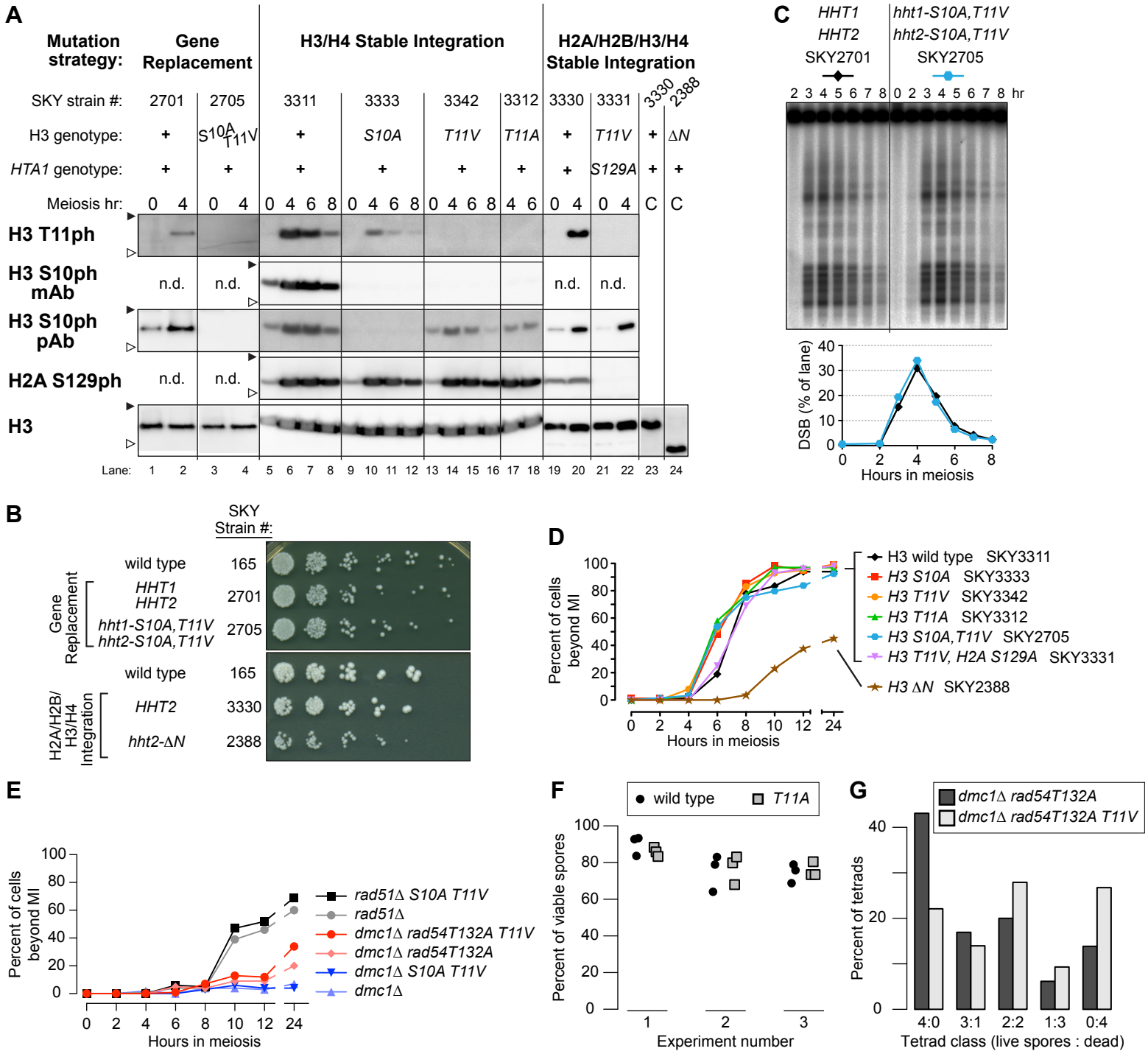
Kniewel et al., Figure 2



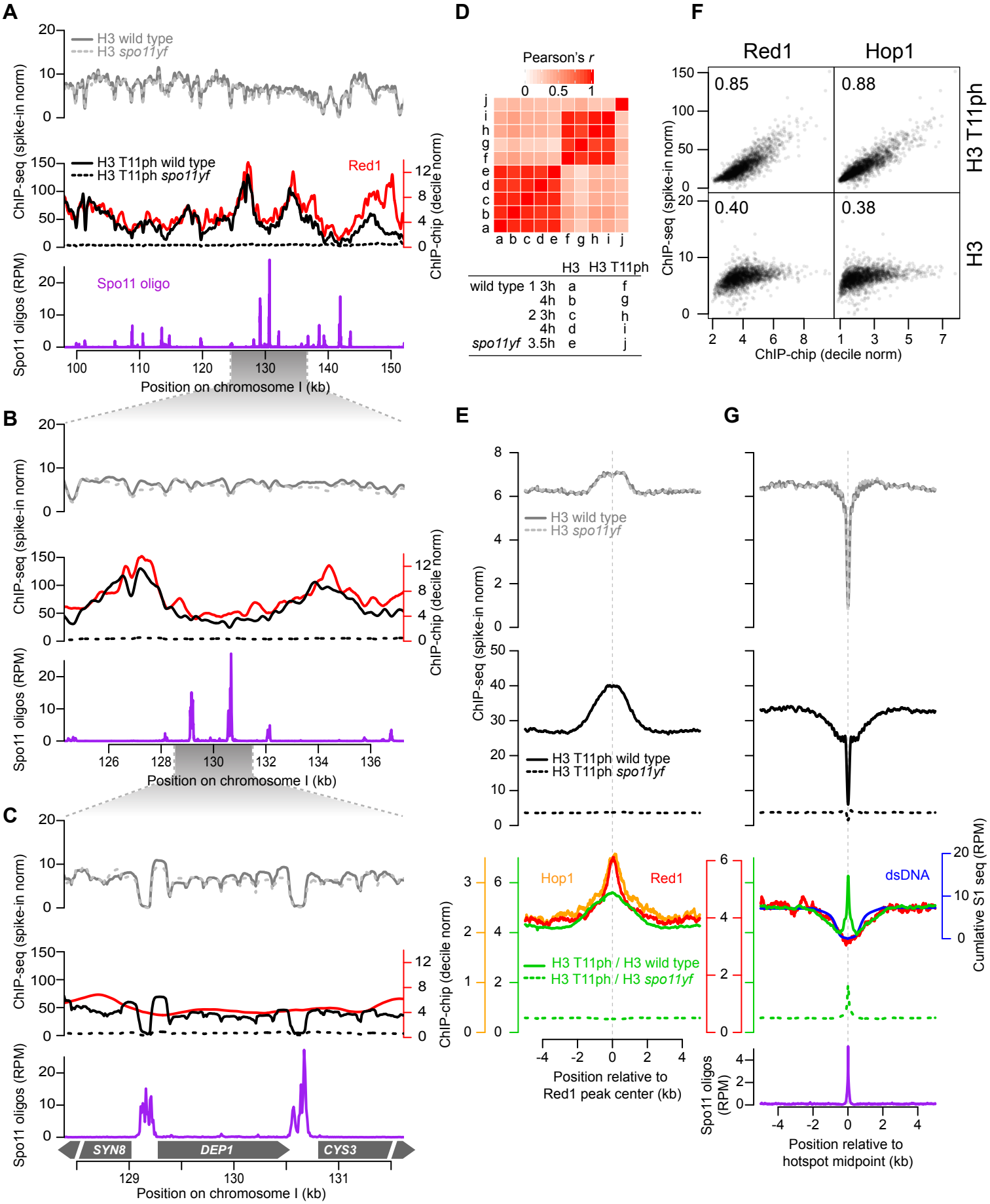
Kniewel et al., Figure 3



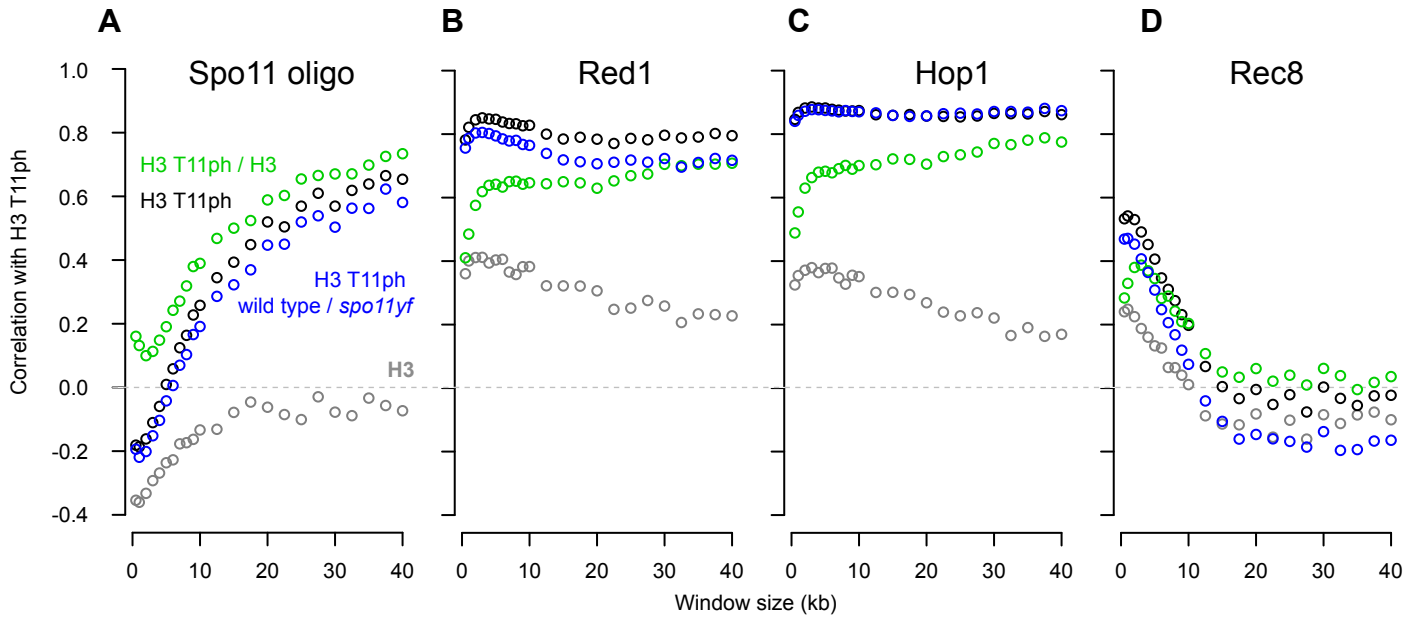
Kniewel et al., Figure 4



Kniewel et al., Figure 5

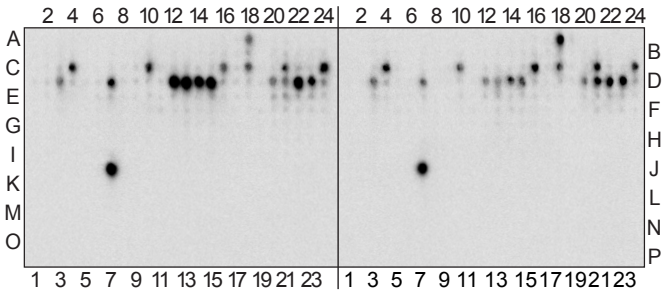


Kniewel et al., Figure 6



Kniewel et al., Supplemental Figure S1

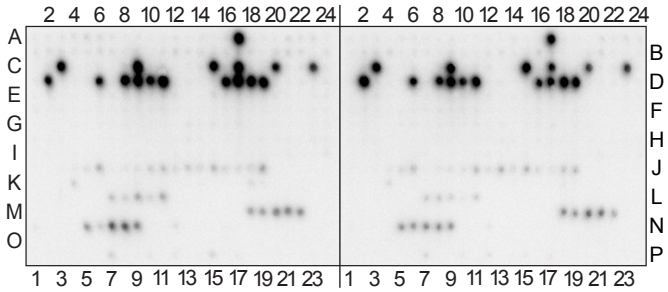
A **(i) H3 T11ph mAb**



Highly Positive Peptides:

A18	H3 1-19 **T11ph**	D12	H3 1-19 R8me2s K9me1 **T11ph**
C4	H3 1-19 R8me2s **T11ph**	D13	H3 1-19 R8me2s K9me2 **T11ph**
C10	H3 1-19 R8me2a **T11ph**	D14	H3 1-19 R8me2s K9me2 **T11ph**
C16	H3 1-19 R8cit **T11ph**	D15	H3 1-19 R8me2s K9ac **T11ph**
C18	H3 1-19 K9me1 **T11ph**	D20	H3 1-19 R8me2a K9me1 **T11ph**
C21	H3 1-19 K9me2 **T11ph**	D21	H3 1-19 R8me2a K9me2 **T11ph**
C24	H3 1-19 K9me3 **T11ph**	D22	H3 1-19 R8me2a K9me3 **T11ph**
D3	H3 1-19 K9ac **T11ph**	D23	H3 1-19 R8me2s K9ac **T11ph**
D7	H3 1-19 **T11ph** K14ac	J7	H3 7-26 **T11ph** K14ac

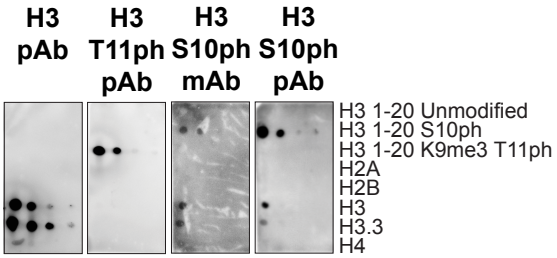
(ii) H3 S10ph pAb



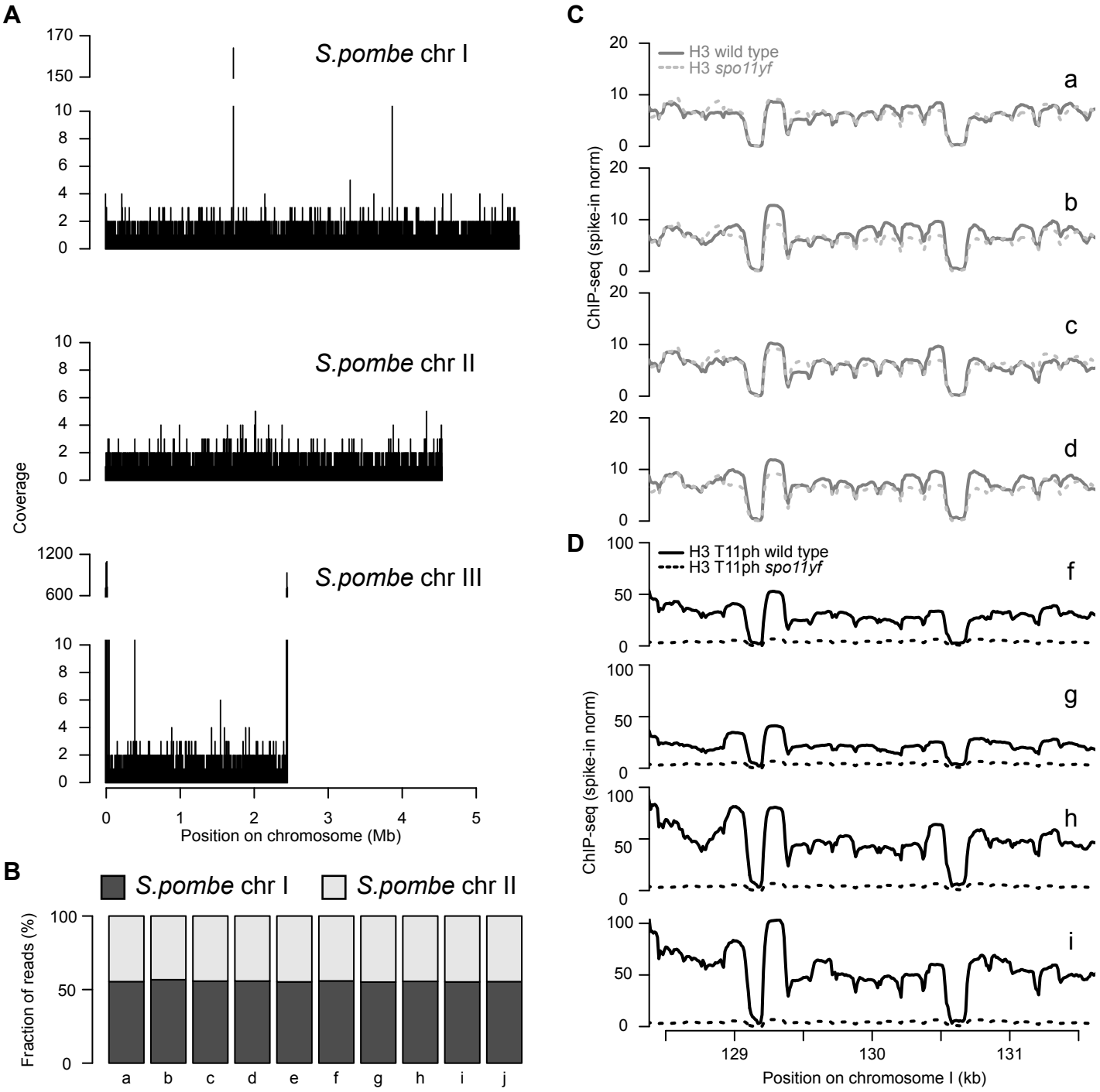
Highly Positive Peptides:

A17	H3 1-19 **S10ph**	D8	H3 1-19 R8me2s K9me1 **S10ph**
C3	H3 1-19 R8me2s **S10ph**	D9	H3 1-19 R8me2s K9me2 **S10ph**
C9	H3 1-19 R8me2a **S10ph**	D10	H3 1-19 R8me2s K9me2 **S10ph**
C15	H3 1-19 R8cit **S10ph**	D11	H3 1-19 R8me2s K9ac **S10ph**
C17	H3 1-19 K9me1 **S10ph**	D16	H3 1-19 R8me2a K9me1 **S10ph**
C20	H3 1-19 K9me2 **S10ph**	D17	H3 1-19 R8me2a K9me2 **S10ph**
C23	H3 1-19 K9me3 **S10ph**	D18	H3 1-19 R8me2a K9me3 **S10ph**
D2	H3 1-19 K9ac **S10ph**	D19	H3 1-19 R8me2s K9ac **S10ph**
D6	H3 1-19 **S10ph** K14ac		

B



Kniewel et al., Supplemental Figure S2



Supplemental Table S1. List of *S. cerevisiae* and *S. pombe* strains used in this study.

***S. cerevisiae*, SK1 background[a]**		
Strain SKY#	**Genotype**	**Reference or source**
Strains for Figure 1: Genetic requirements of H3 T11ph		
165[a]		(CHA *et al.* 2000)
198	*HO, lys2, ura3::hisG, spo11-Y135F-HA::URA*	(CHA *et al.* 2000)
50	*leu2, arg4-Nsp, nuc1Δ::LEU2, rad50-K81I (rad50S)::URA3*	(LIU *et al.* 1995)
2051	*ndt80Δ::LEU2*	(XU *et al.* 1995)
3455	*ho::hisG, ura3(Δsma-pst), rad51Δ::hisG-URA3-hisG*	Neil Hunter
2578	*his4-X, dmc1Δ::LEU2*	(BISHOP *et al.* 1992)
Strain for Figure 3: H3 T11ph kinase determination		
3095	*his4-X/his4-B, ura3::GST-mek1-as1::URA3/ura3, mek1Δ::kanMX6, dmc1Δ::LEU2*	(WAN *et al.* 2004)
Strains for Table 1, Table 2 and Figure 4: H3 T11 mutants		
2701	*HHT1::kanMX4, HHT2::hphMX4*	This study
2705	*hht1-S10A, T11V::kanMX4; hht2-S10A, T11V::hphMX4*	This study
3166	***MATa**, ho::LYS2, lys2, leu2::hisG, ura3, hht1-hhf1Δ::kanMX, hhf2-hht2Δ::natMX, hta2-htb2Δ::natMX, pRK12[CEN6/ARS4, URA3, HTA1-HTB1, HHF2-HHT2]*	This study
3167	***MATα**, ho::LYS2, lys2, leu2::hisG, ura3, hht1-hhf1Δ::kanMX, hhf2-hht2Δ::natMX, hta2-htb2Δ::natMX, pRK12[CEN6/ARS4, URA3, HTA1-HTB1, HHF2-HHT2]*	This study
3311	*hht1-hhf1Δ::kanMX, hhf2-hht2Δ::natMX, hta2-htb2Δ::natMX, leu2::HHF2-HHT2::LEU2*	This study
3333	*hht1-hhf1Δ::kanMX, hhf2-hht2Δ::natMX, hta2-htb2Δ::natMX, leu2::HHF2-hht2-S10A::LEU2*	This study
3342	*hht1-hhf1Δ::kanMX, hhf2-hht2Δ::natMX, hta2-htb2Δ::natMX, leu2::HHF2-hht2-T11V::LEU2*	This study
3334	*hht1-hhf1Δ::kanMX, hhf2-hht2Δ::natMX, hta2-htb2Δ::natMX; leu2::HHF2-hht2-S10A, T11V::LEU2*	This study
3312	*hht1-hhf1Δ::kanMX, hhf2-hht2Δ::natMX, hta2-htb2Δ::natMX, leu2::HHF2-hht2-T11A::LEU2*	This study
3313	*hht1-hhf1Δ::kanMX, hhf2-hht2Δ::natMX, hta2-htb2Δ::natMX, leu2::HHF2-hht2-T11S::LEU2*	This study

Supplemental Table 1 (continued).

3332	*hht1-hhf1Δ::kanMX, hhf2-hht2Δ::natMX, hta2-htb2Δ::natMX, leu2::HHF2-hht2-T11D::LEU2*	This study
3303	*hht1-hhf1Δ::kanMX, hhf2-hht2Δ::natMX, hta2-htb2Δ::natMX, leu2::HHF2-hht2-T11E::LEU2*	This study
2283	*hht1-hhf1Δ::kanMX, hhf2-hht2Δ::natMX, hta2-htb2Δ::natMX, hta1-htb1Δ::hphMX, pRK12[CEN6/ARS4, URA3, HTA1-HTB1, HHF2-HHT2]*	This study
3330	*hht1-hhf1Δ::kanMX, hhf2-hht2Δ::natMX, hta2-htb2Δ::natMX, hta1-htb1Δ::hphMX, leu2::HTA1-HTB1-HHF2-HHT2-LEU2*	This study
3264	*hht1-hhf1Δ::kanMX, hhf2-hht2Δ::natMX, hta2-htb2Δ::natMX, hta1-htb1Δ::hphMX, leu2::HTA1-HTB1-HHF2-hht2-T11V::LEU2*	This study
2388	*hht1-hhf1Δ::kanMX, hhf2-hht2Δ::natMX, hta2-htb2Δ::natMX, hta1-htb1Δ::hphMX, leu2::HTA1-HTB1-HHF2-hht2-Δ1-30(ΔN)::LEU2*	This study
3428	*hht1-hhf1Δ::kanMX, hhf2-hht2Δ::natMX, hta2-htb2Δ::natMX, leu2::HHF2-HHT2::LEU2, arg4-Nsp/arg4-Bgl*	This study & (MARTINI *et al.* 2006)
3431	*hht1-hhf1Δ::kanMX, hhf2-hht2Δ::natMX, hta2-htb2Δ::natMX, leu2::HHF2-hht2-T11A::LEU2, arg4-Nsp/arg4-Bgl*	This study & (MARTINI *et al.* 2006)
3438, 3439, 3440[b]	*trp1::hisG, his4-N/his4-G, hhf1-hht1Δ::LEU2, hhf2-hht2Δ::trp1::kanMX3, pRK92[CEN, ARS, TRP1, HHT2-HHF2]*	(GOVIN *et al.* 2010)
3441, 3442, 3443[b]	*trp1::hisG, his4-N/his4-G, hhf1-hht1Δ::LEU2, hhf2-hht2Δ::trp1::kanMX3, pRK93[CEN, ARS, TRP1, hht2-T11A-HHF2]*	(GOVIN *et al.* 2010)
3265	*hht1-hhf1Δ::kanMX, hhf2-hht2Δ::natMX, hta2-htb2Δ::natMX, hta1-htb1Δ::hphMX, leu2::hta1-S129A-HTB1-HHT2-HHF2::LEU2*	This study
3331	*hht1-hhf1Δ::kanMX, hhf2-hht2Δ::natMX, hta2-htb2Δ::natMX, hta1-htb1Δ::hphMX, leu2::hta1-S129A-HTB1-HHF2-hht2-T11V::LEU2*	This study

Supplemental Table 1 (continued).

4415	*leu2::hisG or leu2-K, arg4-nsp, bgl or ARG4, HHT1::kanMX4, HHT2::hphMX4, set1Δ::kanMX*	This study & ORT4784 X ORT 4785 (SOLLIER *et al.* 2004)
3329	*leu2::hisG or leu2-K, arg4-nsp, bgl or ARG4, hht1-S10A, T11V::kanMX4, hht2-S10A, T11V::hphMX4, set1Δ::kanMX*	This study
3183	*HHT1::kanMX/", HHT2::hygMX/", rad51Δ::hisG-URA3-hisG/"*	This study
3186	*hht1-S10AT11V::kanMX/", hht2-S10AT11V::hygMX/", rad51Δ::hisG-URA3-hisG/"*	This study
3802	*his4X, ura3::RAD54-T132A::URA3, dmc1Δ::hphMX4, rad54::kanMX6*	(NIU *et al.* 2009)
3659	*HIS4, ura3::RAD54-T132A::URA3, dmc1Δ::hphMX4, rad54::kanMX6, hht1-hhf1Δ::kanMX, hhf2-hht2Δ::natMX, hta2-htb2Δ::natMX, leu2::HHF2-hht2-T11V::LEU2*	This study
3078	*HHT1::kanMX4, HHT2::hphMX4, dmc1Δ::LEU2*	This study
3091	*hht1-S10A, T11V::kanMX4, hht2-S10A, T11V::hphMX4, dmc1Δ::LEU2*	This study

Strains for Figure 2: *S. pombe*, Standard background		
2594	*h+, pat1-114, ade6-3049*	(STEINER AND SMITH 2005)
2595	*h+, pat1-114, ade6-3049, rad50-K81I (rad50S)*	(YOUNG *et al.* 2002)
2596	*h-, pat1-114, ade6-3049, ura4-DIB, rec12-171::ura4*$^{+}$	(DAVIS AND SMITH 2003)

[a] All *S. cerevisiae* strains are diploid *MAT***a***/MATα, ho::LYS2/", lys2/", leu2::hisG/", ura3/"* (except SKY3166 and SKY3167) and homozygous at all loci unless otherwise noted (KANE AND ROTH 1974).
[b] Three independent plasmid shuffle transformants.

Supplemental Table 1 (continued).

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Supplemental Table S2. Synthetic peptide array.

Active Motif
MODified™ Histone Peptide Array*
Catalog Nos. 13001 & 13005

Modification number	*Peptide location*	*Peptide sequence*	*name*	*Mod1*	*Mod2*	*Mod 3*	*Mod 4*	*N-terminus*
1	A 1	A R T K Q T A R K S T G G K A P R K Q	H3 1-19	unmod				free
2	A 2	A Rme2s T K Q T A R K S T G G K A P R K Q	H3 1-19	R2me2s				free
3	A 3	A Rme2a T K Q T A R K S T G G K A P R K Q	H3 1-19	R2me2a				free
4	A 4	A Cit T K Q T A R K S T G G K A P R K Q	H3 1-19	R2Citr				free
5	A 5	A R pT K Q T A R K S T G G K A P R K Q	H3 1-19	T3P				free
6	A 6	A R T Kme1 Q T A R K S T G G K A P R K Q	H3 1-19	K4me1				free
7	A 7	A R T Kme2 Q T A R K S T G G K A P R K Q	H3 1-19	K4me2				free
8	A 8	A R T Kme3 Q T A R K S T G G K A P R K Q	H3 1-19	K4me3				free
9	A 9	A R T Kac Q T A R K S T G G K A P R K Q	H3 1-19	K4ac				free
10	A10	A R T K Q T A Rme2s K S T G G K A P R K Q	H3 1-19	R8me2s				free
11	A11	A R T K Q T A Rme2a K S T G G K A P R K Q	H3 1-19	R8me2a				free
12	A12	A R T K Q T A Cit K S T G G K A P R K Q	H3 1-19	R8Citr				free
13	A13	A R T K Q T A R Kme1 S T G G K A P R K Q	H3 1-19	K9me				free
14	A14	A R T K Q T A R Kme2 S T G G K A P R K Q	H3 1-19	K9m2				free
15	A15	A R T K Q T A R Kme3 S T G G K A P R K Q	H3 1-19	K9me3				free
16	A16	A R T K Q T A R Kac S T G G K A P R K Q	H3 1-19	K9ac				free
17	A17	A R T K Q T A R K pS T G G K A P R K Q	H3 1-19	S10P				free
18	A18	A R T K Q T A R K S pT G G K A P R K Q	H3 1-19	T11P				free
19	A19	A R T K Q T A R K S T G G Kac A P R K Q	H3 1-19	K14ac				free
20	A20	A Rme2s pT K Q T A R K S T G G K A P R K Q	H3 1-19	R2me2s	T3P			free
21	A21	A Rme2s T Kme1 Q T A R K S T G G K A P R K Q	H3 1-19	R2me2s	K4me1			free
22	A22	A Rme2s T Kme2 Q T A R K S T G G K A P R K Q	H3 1-19	R2me2s	K4me2			free
23	A23	A Rme2s T Kme3 Q T A R K S T G G K A P R K Q	H3 1-19	R2me2s	K4me3			free
24	A24	A Rme2s T Kac Q T A R K S T G G K A P R K Q	H3 1-19	R2me2s	K4ac			free
25	B 1	A Rme2a pT K Q T A R K S T G G K A P R K Q	H3 1-19	R2me2a	T3P			free
26	B 2	A Rme2a T Kme1 Q T A R K S T G G K A P R K Q	H3 1-19	R2me2a	K4me1			free
27	B 3	A Rme2a T Kme2 Q T A R K S T G G K A P R K Q	H3 1-19	R2me2a	K4me2			free
28	B 4	A Rme2a T Kme3 Q T A R K S T G G K A P R K Q	H3 1-19	R2me2a	K4me3			free
29	B 5	A Rme2a T Kac Q T A R K S T G G K A P R K Q	H3 1-19	R2me2a	K4ac			free
30	B 6	A Cit pT K Q T A R K S T G G K A P R K Q	H3 1-19	R2Citr	T3P			free
31	B 7	A Cit T Kme1 Q T A R K S T G G K A P R K Q	H3 1-19	R2Citr	K4me1			free
32	B 8	A Cit T Kme2 Q T A R K S T G G K A P R K Q	H3 1-19	R2Citr	K4me2			free
33	B 9	A Cit T Kme3 Q T A R K S T G G K A P R K Q	H3 1-19	R2Citr	K4me3			free
34	B10	A Cit T Kac Q T A R K S T G G K A P R K Q	H3 1-19	R2Citr	K4ac			free
35	B11	A R pT Kme1 Q T A R K S T G G K A P R K Q	H3 1-19	T3P	K4me1			free
36	B12	A R pT Kme2 Q T A R K S T G G K A P R K Q	H3 1-19	T3P	K4me2			free
37	B13	A R pT Kme3 Q T A R K S T G G K A P R K Q	H3 1-19	T3P	K4me3			free
38	B14	A R pT Kac Q T A R K S T G G K A P R K Q	H3 1-19	T3P	K4ac			free
39	B15	A Rme2s pT Kme1 Q T A R K S T G G K A P R K Q	H3 1-19	R2me2s	T3P	K4me1		free
40	B16	A Rme2s pT Kme2 Q T A R K S T G G K A P R K Q	H3 1-19	R2me2s	T3P	K4me2		free
41	B17	A Rme2s pT Kme3 Q T A R K S T G G K A P R K Q	H3 1-19	R2me2s	T3P	K4me3		free
42	B18	A Rme2s pT Kac Q T A R K S T G G K A P R K Q	H3 1-19	R2me2s	T3P	K4ac		free
43	B19	A Rme2a pT Kme1 Q T A R K S T G G K A P R K Q	H3 1-19	R2me2a	T3P	K4me1		free
44	B20	A Rme2a pT Kme2 Q T A R K S T G G K A P R K Q	H3 1-19	R2me2a	T3P	K4me2		free
45	B21	A Rme2a pT Kme3 Q T A R K S T G G K A P R K Q	H3 1-19	R2me2a	T3P	K4me3		free
46	B22	A Rme2a pT Kac Q T A R K S T G G K A P R K Q	H3 1-19	R2me2a	T3P	K4ac		free
47	B23	A R T K Q T A Rme2a Kme1 S T G G K A P R K Q	H3 1-19	R8me2s	K9me			free
48	B24	A R T K Q T A Rme2a Kme2 S T G G K A P R K Q	H3 1-19	R8me2s	K9m2			free
49	C 1	A R T K Q T A Rme2a Kme3 S T G G K A P R K Q	H3 1-19	R8me2s	K9me3			free
50	C 2	A R T K Q T A Rme2a Kac S T G G K A P R K Q	H3 1-19	R8me2s	K9ac			free
51	C 3	A R T K Q T A Rme2a K pS T G G K A P R K Q	H3 1-19	R8me2s	S10P			free
52	C 4	A R T K Q T A Rme2a K S pT G G K A P R K Q	H3 1-19	R8me2s	T11P			free
53	C 5	A R T K Q T A Rme2a Kme1 S T G G K A P R K Q	H3 1-19	R8me2a	K9me			free
54	C 6	A R T K Q T A Rme2a Kme2 S T G G K A P R K Q	H3 1-19	R8me2a	K9m2			free
55	C 7	A R T K Q T A Rme2a Kme3 S T G G K A P R K Q	H3 1-19	R8me2a	K9me3			free
56	C 8	A R T K Q T A Rme2a Kac S T G G K A P R K Q	H3 1-19	R8me2a	K9ac			free
57	C 9	A R T K Q T A Rme2a K pS T G G K A P R K Q	H3 1-19	R8me2a	S10P			free
58	C10	A R T K Q T A Rme2a K S pT G G K A P R K Q	H3 1-19	R8me2a	T11P			free
59	C11	A R T K Q T A Cit Kme1 S T G G K A P R K Q	H3 1-19	R8Citr	K9me			free
60	C12	A R T K Q T A Cit Kme2 S T G G K A P R K Q	H3 1-19	R8Citr	K9m2			free
61	C13	A R T K Q T A Cit Kme3 S T G G K A P R K Q	H3 1-19	R8Citr	K9me3			free
62	C14	A R T K Q T A Cit Kac S T G G K A P R K Q	H3 1-19	R8Citr	K9ac			free
63	C15	A R T K Q T A Cit K pS T G G K A P R K Q	H3 1-19	R8Citr	S10P			free
64	C16	A R T K Q T A Cit K S pT G G K A P R K Q	H3 1-19	R8Citr	T11P			free
65	C17	A R T K Q T A R Kme1 pS T G G K A P R K Q	H3 1-19	K9me	S10P			free
66	C18	A R T K Q T A R Kme1 S pT G G K A P R K Q	H3 1-19	K9me	T11P			free
67	C19	A R T K Q T A R Kme1 S T G G Kac A P R K Q	H3 1-19	K9me	K14ac			free
68	C20	A R T K Q T A R Kme2 pS T G G K A P R K Q	H3 1-19	K9me2	S10P			free
69	C21	A R T K Q T A R Kme2 S pT G G K A P R K Q	H3 1-19	K9me2	T11P			free
70	C22	A R T K Q T A R Kme2 S T G G Kac A P R K Q	H3 1-19	K9me2	K14ac			free
71	C23	A R T K Q T A R Kme3 pS T G G K A P R K Q	H3 1-19	K9me3	S10P			free
72	C24	A R T K Q T A R Kme3 S pT G G K A P R K Q	H3 1-19	K9me3	T11P			free
73	D 1	A R T K Q T A R Kme3 S T G G Kac A P R K Q	H3 1-19	K9me3	K14ac			free
74	D 2	A R T K Q T A R Kac pS T G G K A P R K Q	H3 1-19	K9ac	S10P			free
75	D 3	A R T K Q T A R Kac S pT G G K A P R K Q	H3 1-19	K9ac	T11P			free
76	D 4	A R T K Q T A R Kac S T G G Kac A P R K Q	H3 1-19	K9ac	K14ac			free
77	D 5	A R T K Q T A R K pS pT G G K A P R K Q	H3 1-19	S10P	T11P			free
78	D 6	A R T K Q T A R K pS T G G Kac A P R K Q	H3 1-19	S10P	K14ac			free
79	D 7	A R T K Q T A R K S pT G G Kac A P R K Q	H3 1-19	T11P	K14ac			free
80	D 8	A R T K Q T A Rme2s Kme1 pS T G G K A P R K Q	H3 1-19	R8me2s	K9me	S10P		free
81	D 9	A R T K Q T A Rme2s Kme2 pS T G G K A P R K Q	H3 1-19	R8me2s	K9me2	S10P		free
82	D10	A R T K Q T A Rme2s Kme3 pS T G G K A P R K Q	H3 1-19	R8me2s	K9me3	S10P		free
83	D11	A R T K Q T A Rme2s Kac pS T G G K A P R K Q	H3 1-19	R8me2s	K9ac	S10P		free
84	D12	A R T K Q T A Rme2s Kme1 S pT G G K A P R K Q	H3 1-19	R8me2s	K9me	T11P		free
85	D13	A R T K Q T A Rme2s Kme2 S pT G G K A P R K Q	H3 1-19	R8me2s	K9me2	T11P		free

86	D14	A R T K Q T A Rme2s Kme3 S pT G G K A P R K Q	H3 1-19	R8me2s	K9me3	T11P		free
87	D15	A R T K Q T A Rme2s Kac S pT G G K A P R K Q	H3 1-19	R8me2s	K9ac	T11P		free
88	D16	A R T K Q T A Rme2a Kme1 pS T G G K A P R K Q	H3 1-19	R8me2a	K9me	S10P		free
89	D17	A R T K Q T A Rme2a Kme2 pS T G G K A P R K Q	H3 1-19	R8me2a	K9me2	S10P		free
90	D18	A R T K Q T A Rme2a Kme3 pS T G G K A P R K Q	H3 1-19	R8me2a	K9me3	S10P		free
91	D19	A R T K Q T A Rme2a Kac pS T G G K A P R K Q	H3 1-19	R8me2a	K9ac	S10P		free
92	D20	A R T K Q T A Rme2a Kme1 S pT G G K A P R K Q	H3 1-19	R8me2a	K9me	T11P		free
93	D21	A R T K Q T A Rme2a Kme2 S pT G G K A P R K Q	H3 1-19	R8me2a	K9me2	T11P		free
94	D22	A R T K Q T A Rme2a Kme3 S pT G G K A P R K Q	H3 1-19	R8me2a	K9me3	T11P		free
95	D23	A R T K Q T A Rme2a Kac S pT G G K A P R K Q	H3 1-19	R8me2a	K9ac	T11P		free
96	D24	A R T K Q T A Rme2a Kme1 pS pT G G K A P R K Q	H3 1-19	R8me2a	K9me	S10P	T11P	free
97	E 1	A R T K Q T A Rme2a Kme2 pS pT G G K A P R K Q	H3 1-19	R8me2a	K9me2	S10P	T11P	free
98	E 2	A R T K Q T A Rme2a Kme3 pS pT G G K A P R K Q	H3 1-19	R8me2a	K9me3	S10P	T11P	free
99	E 3	A R T K Q T A Rme2a Kac pS pT G G K A P R K Q	H3 1-19	R8me2a	K9ac	S10P	T11P	free
100	E 4	A Rme2s T Kme1 Q T A Rme2s K S T G G K A P R K Q	H3 1-19	R2me2s	K4me1	R8me2s		free
101	E 5	A Rme2s T Kme2 Q T A Rme2s K S T G G K A P R K Q	H3 1-19	R2me2s	K4me2	R8me2s		free
102	E 6	A Rme2s T Kme3 Q T A Rme2s K S T G G K A P R K Q	H3 1-19	R2me2s	K4me3	R8me2s		free
103	E 7	A Rme2s T Kac Q T A Rme2s K S T G G K A P R K Q	H3 1-19	R2me2s	K4ac	R8me2s		free
104	E 8	A Rme2a T Kme1 Q T A Rme2a K S T G G K A P R K Q	H3 1-19	R2me2a	K4me1	R8me2a		free
105	E 9	A Rme2a T Kme2 Q T A Rme2a K S T G G K A P R K Q	H3 1-19	R2me2a	K4me2	R8me2a		free
106	E10	A Rme2a T Kme3 Q T A Rme2a K S T G G K A P R K Q	H3 1-19	R2me2a	K4me3	R8me2a		free
107	E11	A Rme2a T Kac Q T A Rme2a K S T G G K A P R K Q	H3 1-19	R2me2a	K4ac	R8me2a		free
108	E12	A Rme2s T Kme1 Q T A R Kme1 S T G G K A P R K Q	H3 1-19	R2me2s	K4me1	K9me		free
109	E13	A Rme2s T Kme2 Q T A R Kme1 S T G G K A P R K Q	H3 1-19	R2me2s	K4me2	K9me		free
110	E14	A Rme2s T Kme3 Q T A R Kme1 S T G G K A P R K Q	H3 1-19	R2me2s	K4me3	K9me		free
111	E15	A Rme2s T Kac Q T A R Kme1 S T G G K A P R K Q	H3 1-19	R2me2s	K4ac	K9me		free
112	E16	A Rme2a T Kme1 Q T A R Kme2 S T G G K A P R K Q	H3 1-19	R2me2a	K4me1	K9me2		free
113	E17	A Rme2a T Kme2 Q T A R Kme2 S T G G K A P R K Q	H3 1-19	R2me2a	K4me2	K9me2		free
114	E18	A Rme2a T Kme3 Q T A R Kme2 S T G G K A P R K Q	H3 1-19	R2me2a	K4me3	K9me2		free
115	E19	A Rme2a T Kac Q T A R Kme2 S T G G K A P R K Q	H3 1-19	R2me2a	K4ac	K9me2		free
116	E20	A Rme2s T Kme1 Q T A R Kme3 S T G G K A P R K Q	H3 1-19	R2me2s	K4me1	K9me3		free
117	E21	A Rme2s T Kme2 Q T A R Kme3 S T G G K A P R K Q	H3 1-19	R2me2s	K4me2	K9me3		free
118	E22	A Rme2s T Kme3 Q T A R Kme3 S T G G K A P R K Q	H3 1-19	R2me2s	K4me3	K9me3		free
119	E23	A Rme2s T Kac Q T A R Kme3 S T G G K A P R K Q	H3 1-19	R2me2s	K4ac	K9me3		free
120	E24	A Rme2a T Kme1 Q T A R Kac S T G G K A P R K Q	H3 1-19	R2me2a	K4me1	K9ac		free
121	F 1	A Rme2a T Kme2 Q T A R Kac S T G G K A P R K Q	H3 1-19	R2me2a	K4me2	K9ac		free
122	F 2	A Rme2a T Kme3 Q T A R Kac S T G G K A P R K Q	H3 1-19	R2me2a	K4me3	K9ac		free
123	F 3	A Rme2a T Kac Q T A R Kac S T G G K A P R K Q	H3 1-19	R2me2a	K4ac	K9ac		free
124	F 4	A R T Kme1 Q T A Rme2s Kme1 S T G G K A P R K Q	H3 1-19	K4me1	R8me2s	K9me		free
125	F 5	A R T Kme2 Q T A Rme2s Kme1 S T G G K A P R K Q	H3 1-19	K4me2	R8me2s	K9me		free
126	F 6	A R T Kme3 Q T A Rme2s Kme1 S T G G K A P R K Q	H3 1-19	K4me3	R8me2s	K9me		free
127	F 7	A R T Kac Q T A Rme2s Kme1 S T G G K A P R K Q	H3 1-19	K4ac	R8me2s	K9me		free
128	F 8	A R T Kme1 Q T A Rme2a Kme1 S T G G K A P R K Q	H3 1-19	K4me1	R8me2a	K9me		free
129	F 9	A R T Kme2 Q T A Rme2a Kme1 S T G G K A P R K Q	H3 1-19	K4me2	R8me2a	K9me		free
130	F10	A R T Kme3 Q T A Rme2a Kme1 S T G G K A P R K Q	H3 1-19	K4me3	R8me2a	K9me		free
131	F11	A R T Kac Q T A Rme2a Kme1 S T G G K A P R K Q	H3 1-19	K4ac	R8me2a	K9me		free
132	F12	A R T Kme1 Q T A Rme2s Kme2 S T G G K A P R K Q	H3 1-19	K4me1	R8me2s	K9me2		free
133	F13	A R T Kme2 Q T A Rme2s Kme2 S T G G K A P R K Q	H3 1-19	K4me2	R8me2s	K9me2		free
134	F14	A R T Kme3 Q T A Rme2s Kme2 S T G G K A P R K Q	H3 1-19	K4me3	R8me2s	K9me2		free
135	F15	A R T Kac Q T A Rme2s Kme2 S T G G K A P R K Q	H3 1-19	K4ac	R8me2s	K9me2		free
136	F16	A R T Kme1 Q T A Rme2a Kme2 S T G G K A P R K Q	H3 1-19	K4me1	R8me2a	K9me2		free
137	F17	A R T Kme2 Q T A Rme2a Kme2 S T G G K A P R K Q	H3 1-19	K4me2	R8me2a	K9me2		free
138	F18	A R T Kme3 Q T A Rme2a Kme2 S T G G K A P R K Q	H3 1-19	K4me3	R8me2a	K9me2		free
139	F19	A R T Kac Q T A Rme2a Kme2 S T G G K A P R K Q	H3 1-19	K4ac	R8me2a	K9me2		free
140	F20	A R T Kme1 Q T A Rme2s Kme3 S T G G K A P R K Q	H3 1-19	K4me1	R8me2s	K9me3		free
141	F21	A R T Kme2 Q T A Rme2s Kme3 S T G G K A P R K Q	H3 1-19	K4me2	R8me2s	K9me3		free
142	F22	A R T Kme3 Q T A Rme2s Kme3 S T G G K A P R K Q	H3 1-19	K4me3	R8me2s	K9me3		free
143	F23	A R T Kac Q T A Rme2s Kme3 S T G G K A P R K Q	H3 1-19	K4ac	R8me2s	K9me3		free
144	F24	A R T Kme1 Q T A Rme2a Kme3 S T G G K A P R K Q	H3 1-19	K4me1	R8me2a	K9me3		free
145	G 1	A R T Kme2 Q T A Rme2a Kme3 S T G G K A P R K Q	H3 1-19	K4me2	R8me2a	K9me3		free
146	G 2	A R T Kme3 Q T A Rme2a Kme3 S T G G K A P R K Q	H3 1-19	K4me3	R8me2a	K9me3		free
147	G 3	A R T Kac Q T A Rme2a Kme3 S T G G K A P R K Q	H3 1-19	K4ac	R8me2a	K9me3		free
148	G 4	A R T Kme1 Q T A Rme2s Kac S T G G K A P R K Q	H3 1-19	K4me1	R8me2s	K9ac		free
149	G 5	A R T Kme2 Q T A Rme2s Kac S T G G K A P R K Q	H3 1-19	K4me2	R8me2s	K9ac		free
150	G 6	A R T Kme3 Q T A Rme2s Kac S T G G K A P R K Q	H3 1-19	K4me3	R8me2s	K9ac		free
151	G 7	A R T Kac Q T A Rme2s Kac S T G G K A P R K Q	H3 1-19	K4ac	R8me2s	K9ac		free
152	G 8	A R T Kme1 Q T A Rme2a Kac S T G G K A P R K Q	H3 1-19	K4me1	R8me2a	K9ac		free
153	G 9	A R T Kme2 Q T A Rme2a Kac S T G G K A P R K Q	H3 1-19	K4me2	R8me2a	K9ac		free
154	G10	A R T Kme3 Q T A Rme2a Kac S T G G K A P R K Q	H3 1-19	K4me3	R8me2a	K9ac		free
155	G11	A R T Kac Q T A Rme2a Kac S T G G K A P R K Q	H3 1-19	K4ac	R8me2a	K9ac		free
156	G12	A Rme2s T Kme1 Q T A Rme2s Kme1 S T G G K A P R K Q	H3 1-19	R2me2s	K4me1	R8me2s	K9me	free
157	G13	A Rme2s T Kme2 Q T A Rme2s Kme1 S T G G K A P R K Q	H3 1-19	R2me2s	K4me2	R8me2s	K9me	free
158	G14	A Rme2s T Kme3 Q T A Rme2s Kme1 S T G G K A P R K Q	H3 1-19	R2me2s	K4me3	R8me2s	K9me	free
159	G15	A Rme2s T Kac Q T A Rme2s Kme1 S T G G K A P R K Q	H3 1-19	R2me2s	K4ac	R8me2s	K9me	free
160	G16	A Rme2a T Kme1 Q T A Rme2s Kme1 S T G G K A P R K Q	H3 1-19	R2me2a	K4me1	R8me2s	K9me	free
161	G17	A Rme2a T Kme2 Q T A Rme2s Kme1 S T G G K A P R K Q	H3 1-19	R2me2a	K4me2	R8me2s	K9me	free
162	G18	A Rme2a T Kme3 Q T A Rme2s Kme1 S T G G K A P R K Q	H3 1-19	R2me2a	K4me3	R8me2s	K9me	free
163	G19	A Rme2a T Kac Q T A Rme2s Kme1 S T G G K A P R K Q	H3 1-19	R2me2a	K4ac	R8me2s	K9me	free
164	G20	A Rme2s T Kme1 Q T A Rme2s Kme2 S T G G K A P R K Q	H3 1-19	R2me2s	K4me1	R8me2s	K9me2	free
165	G21	A Rme2s T Kme2 Q T A Rme2s Kme2 S T G G K A P R K Q	H3 1-19	R2me2s	K4me2	R8me2s	K9me2	free
166	G22	A Rme2s T Kme3 Q T A Rme2s Kme2 S T G G K A P R K Q	H3 1-19	R2me2s	K4me3	R8me2s	K9me2	free
167	G23	A Rme2s T Kac Q T A Rme2s Kme2 S T G G K A P R K Q	H3 1-19	R2me2s	K4ac	R8me2s	K9me2	free
168	G24	A Rme2a T Kme1 Q T A Rme2s Kme2 S T G G K A P R K Q	H3 1-19	R2me2a	K4me1	R8me2s	K9me2	free
169	H 1	A Rme2a T Kme2 Q T A Rme2s Kme2 S T G G K A P R K Q	H3 1-19	R2me2a	K4me2	R8me2s	K9me2	free
170	H 2	A Rme2a T Kme3 Q T A Rme2s Kme2 S T G G K A P R K Q	H3 1-19	R2me2a	K4me3	R8me2s	K9me2	free
171	H 3	A Rme2a T Kac Q T A Rme2s Kme2 S T G G K A P R K Q	H3 1-19	R2me2a	K4ac	R8me2s	K9me2	free
172	H 4	A Rme2s T Kme1 Q T A Rme2s Kme3 S T G G K A P R K Q	H3 1-19	R2me2s	K4me1	R8me2s	K9me3	free
173	H 5	A Rme2s T Kme2 Q T A Rme2s Kme3 S T G G K A P R K Q	H3 1-19	R2me2s	K4me2	R8me2s	K9me3	free
174	H 6	A Rme2s T Kme3 Q T A Rme2s Kme3 S T G G K A P R K Q	H3 1-19	R2me2s	K4me3	R8me2s	K9me3	free
175	H 7	A Rme2s T Kac Q T A Rme2s Kme3 S T G G K A P R K Q	H3 1-19	R2me2s	K4ac	R8me2s	K9me3	free
176	H 8	A Rme2a T Kme1 Q T A Rme2s Kme3 S T G G K A P R K Q	H3 1-19	R2me2a	K4me1	R8me2s	K9me3	free
177	H 9	A Rme2a T Kme2 Q T A Rme2s Kme3 S T G G K A P R K Q	H3 1-19	R2me2a	K4me2	R8me2s	K9me3	free
178	H10	A Rme2a T Kme3 Q T A Rme2s Kme3 S T G G K A P R K Q	H3 1-19	R2me2a	K4me3	R8me2s	K9me3	free
179	H11	A Rme2a T Kac Q T A Rme2s Kme3 S T G G K A P R K Q	H3 1-19	R2me2a	K4ac	R8me2s	K9me3	free

180	H12	A Rme2s T Kme1 Q T A Rme2s Kac S T G G K A P R K Q	H3 1-19	R2me2s	K4me1	R8me2s	K9ac	free
181	H13	A Rme2s T Kme2 Q T A Rme2s Kac S T G G K A P R K Q	H3 1-19	R2me2s	K4me2	R8me2s	K9ac	free
182	H14	A Rme2s T Kme3 Q T A Rme2s Kac S T G G K A P R K Q	H3 1-19	R2me2s	K4me3	R8me2s	K9ac	free
183	H15	A Rme2s T Kac Q T A Rme2s Kac S T G G K A P R K Q	H3 1-19	R2me2s	K4ac	R8me2s	K9ac	free
184	H16	A Rme2a T Kme1 Q T A Rme2s Kac S T G G K A P R K Q	H3 1-19	R2me2a	K4me1	R8me2s	K9ac	free
185	H17	A Rme2a T Kme2 Q T A Rme2s Kac S T G G K A P R K Q	H3 1-19	R2me2a	K4me2	R8me2s	K9ac	free
186	H18	A Rme2a T Kme3 Q T A Rme2s Kac S T G G K A P R K Q	H3 1-19	R2me2a	K4me3	R8me2s	K9ac	free
187	H19	A Rme2a T Kac Q T A Rme2s Kac S T G G K A P R K Q	H3 1-19	R2me2a	K4ac	R8me2s	K9ac	free
188	H20	A Rme2s T Kme1 Q T A Rme2a Kme1 S T G G K A P R K Q	H3 1-19	R2me2s	K4me1	R8me2a	K9me	free
189	H21	A Rme2s T Kme2 Q T A Rme2a Kme1 S T G G K A P R K Q	H3 1-19	R2me2s	K4me2	R8me2a	K9me	free
190	H22	A Rme2s T Kme3 Q T A Rme2a Kme1 S T G G K A P R K Q	H3 1-19	R2me2s	K4me3	R8me2a	K9me	free
191	H23	A Rme2s T Kac Q T A Rme2a Kme1 S T G G K A P R K Q	H3 1-19	R2me2s	K4ac	R8me2a	K9me	free
192	H24	A Rme2a T Kme1 Q T A Rme2a Kme1 S T G G K A P R K Q	H3 1-19	R2me2a	K4me1	R8me2a	K9me	free
193	I 1	A Rme2a T Kme2 Q T A Rme2a Kme1 S T G G K A P R K Q	H3 1-19	R2me2a	K4me2	R8me2a	K9me	free
194	I 2	A Rme2a T Kme3 Q T A Rme2a Kme1 S T G G K A P R K Q	H3 1-19	R2me2a	K4me3	R8me2a	K9me	free
195	I 3	A Rme2a T Kac Q T A Rme2a Kme1 S T G G K A P R K Q	H3 1-19	R2me2a	K4ac	R8me2a	K9me	free
196	I 4	A Rme2s T Kme1 Q T A Rme2a Kme2 S T G G K A P R K Q	H3 1-19	R2me2s	K4me1	R8me2a	K9me2	free
197	I 5	A Rme2s T Kme2 Q T A Rme2a Kme2 S T G G K A P R K Q	H3 1-19	R2me2s	K4me2	R8me2a	K9me2	free
198	I 6	A Rme2s T Kme3 Q T A Rme2a Kme2 S T G G K A P R K Q	H3 1-19	R2me2s	K4me3	R8me2a	K9me2	free
199	I 7	A Rme2s T Kac Q T A Rme2a Kme2 S T G G K A P R K Q	H3 1-19	R2me2s	K4ac	R8me2a	K9me2	free
200	I 8	A Rme2a T Kme1 Q T A Rme2a Kme2 S T G G K A P R K Q	H3 1-19	R2me2a	K4me1	R8me2a	K9me2	free
201	I 9	A Rme2a T Kme2 Q T A Rme2a Kme2 S T G G K A P R K Q	H3 1-19	R2me2a	K4me2	R8me2a	K9me2	free
202	I10	A Rme2a T Kme3 Q T A Rme2a Kme2 S T G G K A P R K Q	H3 1-19	R2me2a	K4me3	R8me2a	K9me2	free
203	I11	A Rme2a T Kac Q T A Rme2a Kme2 S T G G K A P R K Q	H3 1-19	R2me2a	K4ac	R8me2a	K9me2	free
204	I12	A Rme2s T Kme1 Q T A Rme2a Kme3 S T G G K A P R K Q	H3 1-19	R2me2s	K4me1	R8me2a	K9me3	free
205	I13	A Rme2s T Kme2 Q T A Rme2a Kme3 S T G G K A P R K Q	H3 1-19	R2me2s	K4me2	R8me2a	K9me3	free
206	I14	A Rme2s T Kme3 Q T A Rme2a Kme3 S T G G K A P R K Q	H3 1-19	R2me2s	K4me3	R8me2a	K9me3	free
207	I15	A Rme2s T Kac Q T A Rme2a Kme3 S T G G K A P R K Q	H3 1-19	R2me2s	K4ac	R8me2a	K9me3	free
208	I16	A Rme2a T Kme1 Q T A Rme2a Kme3 S T G G K A P R K Q	H3 1-19	R2me2a	K4me1	R8me2a	K9me3	free
209	I17	A Rme2a T Kme2 Q T A Rme2a Kme3 S T G G K A P R K Q	H3 1-19	R2me2a	K4me2	R8me2a	K9me3	free
210	I18	A Rme2a T Kme3 Q T A Rme2a Kme3 S T G G K A P R K Q	H3 1-19	R2me2a	K4me3	R8me2a	K9me3	free
211	I19	A Rme2a T Kac Q T A Rme2a Kme3 S T G G K A P R K Q	H3 1-19	R2me2a	K4ac	R8me2a	K9me3	free
212	I20	A Rme2s T Kme1 Q T A Rme2a Kac S T G G K A P R K Q	H3 1-19	R2me2s	K4me1	R8me2a	K9ac	free
213	I21	A Rme2s T Kme2 Q T A Rme2a Kac S T G G K A P R K Q	H3 1-19	R2me2s	K4me2	R8me2a	K9ac	free
214	I22	A Rme2s T Kme3 Q T A Rme2a Kac S T G G K A P R K Q	H3 1-19	R2me2s	K4me3	R8me2a	K9ac	free
215	I23	A Rme2s T Kac Q T A Rme2a Kac S T G G K A P R K Q	H3 1-19	R2me2s	K4ac	R8me2a	K9ac	free
216	I24	A Rme2a T Kme1 Q T A Rme2a Kac S T G G K A P R K Q	H3 1-19	R2me2a	K4me1	R8me2a	K9ac	free
217	J 1	A Rme2a T Kme2 Q T A Rme2a Kac S T G G K A P R K Q	H3 1-19	R2me2a	K4me2	R8me2a	K9ac	free
218	J 2	A Rme2a T Kme3 Q T A Rme2a Kac S T G G K A P R K Q	H3 1-19	R2me2a	K4me3	R8me2a	K9ac	free
219	J 3	A Rme2a T Kac Q T A Rme2a Kac S T G G K A P R K Q	H3 1-19	R2me2a	K4ac	R8me2a	K9ac	free
220	J 4	A R K S T G G K A P R K Q L A T K A A R	H3 7-26	unmod				acetylated
221	J 5	A R K S T G G Kac A P R K Q L A T K A A R	H3 7-26	K14ac				acetylated
222	J 6	A R K pS T G G Kac A P R K Q L A T K A A R	H3 7-26	K14ac	S10P			acetylated
223	J 7	A R K S pT G G Kac A P R K Q L A T K A A R	H3 7-26	K14ac	T11P			acetylated
224	J 8	A R K S T G G K A P Rme2s K Q L A T K A A R	H3 7-26	R17me2s				acetylated
225	J 9	A R K S T G G K A P Rme2a K Q L A T K A A R	H3 7-26	R17me2a				acetylated
226	J10	A R K S T G G K A P Cit K Q L A T K A A R	H3 7-26	R17Citr				acetylated
227	J11	A R K S T G G K A P R Kac Q L A T K A A R	H3 7-26	K18Ac				acetylated
228	J12	A R K S T G G Kac A P Rme2s K Q L A T K A A R	H3 7-26	K14ac	R17me2s			acetylated
229	J13	A R K S T G G Kac A P Rme2a K Q L A T K A A R	H3 7-26	K14ac	R17me2a			acetylated
230	J14	A R K S T G G Kac A P R Kac Q L A T K A A R	H3 7-26	K14ac	K18Ac			acetylated
231	J15	A R K S T G G K A P Rme2s Kac Q L A T K A A R	H3 7-26	R17me2s	K18Ac			acetylated
232	J16	A R K S T G G K A P Rme2a Kac Q L A T K A A R	H3 7-26	R17me2a	K18Ac			acetylated
233	J17	A R K S T G G K A P Cit Kac Q L A T K A A R	H3 7-26	R17Citr	K18Ac			acetylated
234	J18	A R K S T G G Kac A P Rme2s Kac Q L A T K A A R	H3 7-26	K14ac	R17me2s	K18Ac		acetylated
235	J19	A R K S T G G Kac A P Rme2a Kac Q L A T K A A R	H3 7-26	K14ac	R17me2a	K18Ac		acetylated
236	J20	P R K Q L A T K A A R K S A P A T G G	H3 16-35	unmod				acetylated
237	J21	P R K Q L A T K A A Rme2s K S A P A T G G	H3 16-35	R26me2s				acetylated
238	J22	P R K Q L A T K A A Rme2a K S A P A T G G	H3 16-35	R26me2a				acetylated
239	J23	P R K Q L A T K A A Cit K S A P A T G G	H3 16-35	R26Citr				acetylated
240	J24	P R K Q L A T K A A R Kme1 S A P A T G G	H3 16-35	K27me				acetylated
241	K 1	P R K Q L A T K A A R Kme2 S A P A T G G	H3 16-35	K27me2				acetylated
242	K 2	P R K Q L A T K A A R Kme3 S A P A T G G	H3 16-35	K27me3				acetylated
243	K 3	P R K Q L A T K A A R Kac S A P A T G G	H3 16-35	K27ac				acetylated
244	K 4	P R K Q L A T K A A R K pS A P A T G G	H3 16-35	S28P				acetylated
245	K 5	P R K Q L A T K A A Rme2s Kme1 S A P A T G G	H3 16-35	R26me2s	K27me			acetylated
246	K 6	P R K Q L A T K A A Rme2s Kme2 S A P A T G G	H3 16-35	R26me2s	K27me2			acetylated
247	K 7	P R K Q L A T K A A Rme2s Kme3 S A P A T G G	H3 16-35	R26me2s	K27me3			acetylated
248	K 8	P R K Q L A T K A A Rme2s Kac S A P A T G G	H3 16-35	R26me2s	K27ac			acetylated
249	K 9	P R K Q L A T K A A Rme2s K pS A P A T G G	H3 16-35	R26me2s	S28P			acetylated
250	K10	P R K Q L A T K A A Rme2a Kme1 S A P A T G G	H3 16-35	R26me2a	K27me			acetylated
251	K11	P R K Q L A T K A A Rme2a Kme2 S A P A T G G	H3 16-35	R26me2a	K27me2			acetylated
252	K12	P R K Q L A T K A A Rme2a Kme3 S A P A T G G	H3 16-35	R26me2a	K27me3			acetylated
253	K13	P R K Q L A T K A A Rme2a Kac S A P A T G G	H3 16-35	R26me2a	K27ac			acetylated
254	K14	P R K Q L A T K A A Rme2a K pS A P A T G G	H3 16-35	R26me2a	S28P			acetylated
255	K15	P R K Q L A T K A A Cit Kme1 S A P A T G G	H3 16-35	R26Citr	K27me			acetylated
256	K16	P R K Q L A T K A A Cit Kme2 S A P A T G G	H3 16-35	R26Citr	K27me2			acetylated
257	K17	P R K Q L A T K A A Cit Kme3 S A P A T G G	H3 16-35	R26Citr	K27me3			acetylated
258	K18	P R K Q L A T K A A Cit K pS A P A T G G	H3 16-35	R26Citr	S28P			acetylated
259	K19	P R K Q L A T K A A R Kme1 pS A P A T G G	H3 16-35	K27me	S28P			acetylated
260	K20	P R K Q L A T K A A R Kme2 pS A P A T G G	H3 16-35	K27me2	S28P			acetylated
261	K21	P R K Q L A T K A A R Kme3 pS A P A T G G	H3 16-35	K27me3	S28P			acetylated
262	K22	P R K Q L A T K A A R Kac pS A P A T G G	H3 16-35	K27ac	S28P			acetylated
263	K23	P R K Q L A T K A A Rme2s Kme1 pS A P A T G G	H3 16-35	R26me2s	K27me	S28P		acetylated
264	K24	P R K Q L A T K A A Rme2s Kme2 pS A P A T G G	H3 16-35	R26me2s	K27me2	S28P		acetylated
265	L 1	P R K Q L A T K A A Rme2s Kme3 pS A P A T G G	H3 16-35	R26me2s	K27me3	S28P		acetylated
266	L 2	P R K Q L A T K A A Rme2s Kac pS A P A T G G	H3 16-35	R26me2s	K27ac	S28P		acetylated
267	L 3	P R K Q L A T K A A Rme2a Kme1 pS A P A T G G	H3 16-35	R26me2a	K27me	S28P		acetylated
268	L 4	P R K Q L A T K A A Rme2a Kme2 pS A P A T G G	H3 16-35	R26me2a	K27me2	S28P		acetylated
269	L 5	P R K Q L A T K A A Rme2a Kme3 pS A P A T G G	H3 16-35	R26me2a	K27me3	S28P		acetylated
270	L 6	P R K Q L A T K A A Rme2a Kac pS A P A T G G	H3 16-35	R26me2a	K27ac	S28P		acetylated
271	L 7	R K S A P A T G G V K K P H R Y R P G	H3 26-45	unmod				acetylated
272	L 8	R K S A P A T G G V Kme1 K P H R Y R P G	H3 26-45	K36me				acetylated
273	L 9	R K S A P A T G G V Kme2 K P H R Y R P G	H3 26-45	K36me2				acetylated

274	L10	R K S A P A T G G V Kme3 K P H R Y R P G	H3 26-45	K36me3				acetylated
275	L11	R K S A P A T G G V Kac K P H R Y R P G	H3 26-45	K36ac				acetylated
276	L12	S G R G K G G K G L G K G G A K R H R	H4 1-19	unmod				free
277	L13	pS G R G K G G K G L G K G G A K R H R	H4 1-19	S1P				free
278	L14	S G Rme2s G K G G K G L G K G G A K R H R	H4 1-19	R3me2s				free
279	L15	S G Rme2a G K G G K G L G K G G A K R H R	H4 1-19	R3me2a				free
280	L16	S G R G Kac G G K G L G K G G A K R H R	H4 1-19	K5ac				free
281	L17	S G R G K G G Kac G L G K G G A K R H R	H4 1-19	K8ac				free
282	L18	S G R G K G G K G L G Kac G G A K R H R	H4 1-19	K12ac				free
283	L19	S G R G K G G K G L G K G G A Kac R H R	H4 1-19	K16ac				free
284	L20	pS G Rme2s G K G G K G L G K G G A K R H R	H4 1-19	S1P	R3me2s			free
285	L21	pS G Rme2a G K G G K G L G K G G A K R H R	H4 1-19	S1P	R3me2a			free
286	L22	pS G R G Kac G G K G L G K G G A K R H R	H4 1-19	S1P	K5ac			free
287	L23	S G Rme2s G Kac G G K G L G K G G A K R H R	H4 1-19	R3me2s	K5ac			free
288	L24	S G Rme2s G K G G Kac G L G K G G A K R H R	H4 1-19	R3me2s	K8ac			free
289	M 1	S G Rme2a G Kac G G K G L G K G G A K R H R	H4 1-19	R3me2a	K5ac			free
290	M 2	S G Rme2a G K G G Kac G L G K G G A K R H R	H4 1-19	R3me2a	K8ac			free
291	M 3	S G R G Kac G G Kac G L G K G G A K R H R	H4 1-19	K5ac	K8ac			free
292	M 4	S G R G K G G Kac G L G Kac G G A K R H R	H4 1-19	K8ac	K12ac			free
293	M 5	S G R G K G G Kac G L G K G G A Kac R H R	H4 1-19	K8ac	K16ac			free
294	M 6	S G R G K G G K G L G Kac G G A Kac R H R	H4 1-19	K12ac	K16ac			free
295	M 7	pS G Rme2s G Kac G G K G L G K G G A K R H R	H4 1-19	S1P	R3me2s	K5ac		free
296	M 8	pS G Rme2a G Kac G G K G L G K G G A K R H R	H4 1-19	S1P	R3me2a	K5ac		free
297	M 9	S G Rme2s G Kac G G Kac G L G K G G A K R H R	H4 1-19	R3me2s	K5ac	K8ac		free
298	M10	S G Rme2a G Kac G G Kac G L G K G G A K R H R	H4 1-19	R3me2a	K5ac	K8ac		free
299	M11	S G R G Kac G G Kac G L G Kac G G A K R H R	H4 1-19	K5ac	K8ac	K12ac		free
300	M12	S G R G K G G Kac G L G Kac G G A Kac R H R	H4 1-19	K8ac	K12ac	K16Ac		free
301	M13	pS G Rme2s G Kac G G Kac G L G K G G A K R H R	H4 1-19	S1P	R3me2s	K5ac	K8ac	free
302	M14	pS G Rme2a G Kac G G Kac G L G K G G A K R H R	H4 1-19	S1P	R3me2a	K5ac	K8ac	free
303	M15	S G Rme2s G Kac G G Kac G L G Kac G G A K R H R	H4 1-19	R3me2s	K5ac	K8ac	K12ac	free
304	M16	S G Rme2a G Kac G G Kac G L G Kac G G A K R H R	H4 1-19	R3me2a	K5ac	K8ac	K12ac	free
305	M17	S G R G Kac G G Kac G L G Kac G G A Kac R H R	H4 1-19	K5ac	K8ac	K12ac	K16ac	free
306	M18	G K G G A K R H R K V L R D N I Q G I T	H4 11-30	unmod				acetylated
307	M19	G Kac G G A K R H R K V L R D N I Q G I T	H4 11-30	K12ac				acetylated
308	M20	G K G G A Kac R H R K V L R D N I Q G I T	H4 11-30	K16ac				acetylated
309	M21	G K G G A K Rme2s H R K V L R D N I Q G I T	H4 11-30	R17me2s				acetylated
310	M22	G K G G A K Rme2a H R K V L R D N I Q G I T	H4 11-30	R17me2a				acetylated
311	M23	G K G G A K R H Rme2s K V L R D N I Q G I T	H4 11-30	R19me2s				acetylated
312	M24	G K G G A K R H Rme2a K V L R D N I Q G I T	H4 11-30	R19me2a				acetylated
313	N 1	G K G G A K R H R Kme1 V L R D N I Q G I T	H4 11-30	K20me1				acetylated
314	N 2	G K G G A K R H R Kme2 V L R D N I Q G I T	H4 11-30	K20me2				acetylated
315	N 3	G K G G A K R H R Kme3 V L R D N I Q G I T	H4 11-30	K20me3				acetylated
316	N 4	G K G G A K R H R Kac V L R D N I Q G I T	H4 11-30	K20ac				acetylated
317	N 5	G K G G A K R H R K V L Rme2a D N I Q G I T	H4 11-30	R24me2a				acetylated
318	N 6	G K G G A K R H R K V L Rme2s D N I Q G I T	H4 11-30	R24me2s				acetylated
319	N 7	G Kac G G A Kac R H R K V L R D N I Q G I T	H4 11-30	K12ac	K16ac			acetylated
320	N 8	G K G G A Kac Rme2s H R K V L R D N I Q G I T	H4 11-30	K16ac	R17me2s			acetylated
321	N 9	G K G G A Kac Rme2a H R K V L R D N I Q G I T	H4 11-30	K16ac	R17me2a			acetylated
322	N10	G K G G A Kac R H Rme2s K V L R D N I Q G I T	H4 11-30	K16ac	R19me2s			acetylated
323	N11	G K G G A Kac R H Rme2a K V L R D N I Q G I T	H4 11-30	K16ac	R19me2a			acetylated
324	N12	G K G G A Kac R H R Kme1 V L R D N I Q G I T	H4 11-30	K16ac	K20me1			acetylated
325	N13	G K G G A Kac R H R Kme2 V L R D N I Q G I T	H4 11-30	K16ac	K20me2			acetylated
326	N14	G K G G A Kac R H R Kme3 V L R D N I Q G I T	H4 11-30	K16ac	K20me3			acetylated
327	N15	G K G G A Kac R H R Kac V L R D N I Q G I T	H4 11-30	K16ac	K20ac			acetylated
328	N16	G Kac G G A Kac R H R Kme1 V L R D N I Q G I T	H4 11-30	K12ac	K16ac	K20me1		acetylated
329	N17	G Kac G G A Kac R H R Kme2 V L R D N I Q G I T	H4 11-30	K12ac	K16ac	K20me2		acetylated
330	N18	G Kac G G A Kac R H R Kme3 V L R D N I Q G I T	H4 11-30	K12ac	K16ac	K20me3		acetylated
331	N19	G Kac G G A Kac R H R Kac V L R D N I Q G I T	H4 11-30	K12ac	K16ac	K20ac		acetylated
332	N20	G K G G A K R H Rme2a Kme1 V L R D N I Q G I T	H4 11-30	R19me2a	K20me1			acetylated
333	N21	G K G G A K R H Rme2a Kme2 V L R D N I Q G I T	H4 11-30	R19me2a	K20me2			acetylated
334	N22	G K G G A K R H Rme2a Kme3 V L R D N I Q G I T	H4 11-30	R19me2a	K20me3			acetylated
335	N23	G K G G A K R H Rme2a Kac V L R D N I Q G I T	H4 11-30	R19me2a	K20ac			acetylated
336	N24	G K G G A K R H Rme2s Kme1 V L R D N I Q G I T	H4 11-30	R19me2s	K20me1			acetylated
337	O 1	G K G G A K R H Rme2s Kme2 V L R D N I Q G I T	H4 11-30	R19me2s	K20me2			acetylated
338	O 2	G K G G A K R H Rme2s Kme3 V L R D N I Q G I T	H4 11-30	R19me2s	K20me3			acetylated
339	O 3	G K G G A K R H Rme2s Kac V L R D N I Q G I T	H4 11-30	R19me2s	K20ac			acetylated
340	O 4	G K G G A K R H R Kme1 V L Rme2a D N I Q G I T	H4 11-30	R24me2a	K20me1			acetylated
341	O 5	G K G G A K R H R Kme2 V L Rme2a D N I Q G I T	H4 11-30	R24me2a	K20me2			acetylated
342	O 6	G K G G A K R H R Kme3 V L Rme2a D N I Q G I T	H4 11-30	R24me2a	K20me3			acetylated
343	O 7	G K G G A K R H R Kac V L Rme2a D N I Q G I T	H4 11-30	R24me2a	K20ac			acetylated
344	O 8	G K G G A K R H R Kme1 V L Rme2s D N I Q G I T	H4 11-30	R24me2s	K20me1			acetylated
345	O 9	G K G G A K R H R Kme2 V L Rme2s D N I Q G I T	H4 11-30	R24me2s	K20me2			acetylated
346	O10	G K G G A K R H R Kme3 V L Rme2s D N I Q G I T	H4 11-30	R24me2s	K20me3			acetylated
347	O11	G K G G A K R H R Kac V L Rme2s D N I Q G I T	H4 11-30	R24me2s	K20ac			acetylated
348	O12	S G R G K Q G G K A R A K A K S R S S	H2a 1-19	unmod				free
349	O13	pS G R G K Q G G K A R A K A K S R S S	H2a 1-19	S1P				free
350	O14	S G R G Kac Q G G K A R A K A K S R S S	H2a 1-19	K5ac				free
351	O15	S G R G K Q G G Kac A R A K A K S R S S	H2a 1-19	K9ac				free
352	O16	S G R G K Q G G K A R A Kac A K S R S S	H2a 1-19	K13ac				free
353	O17	pS G R G Kac Q G G K A R A K A K S R S S	H2a 1-19	S1P	K5ac			free
354	O18	pS G R G K Q G G Kac A R A K A K S R S S	H2a 1-19	S1P	K9ac			free
355	O19	pS G R G K Q G G K A R A Kac A K S R S S	H2a 1-19	S1P	K13ac			free
356	O20	S G R G Kac Q G G Kac A R A K A K S R S S	H2a 1-19	K5ac	K9ac			free
357	O21	S G R G Kac Q G G K A R A Kac A K S R S S	H2a 1-19	K5ac	K13ac			free
358	O22	S G R G K Q G G Kac A R A Kac A K S R S S	H2a 1-19	K9ac	K13ac			free
359	O23	pS G R G Kac Q G G Kac A R A K A K S R S S	H2a 1-19	S1P	K5ac	K9ac		free
360	O24	pS G R G Kac Q G G K A R A Kac A K S R S S	H2a 1-19	S1P	K5ac	K13ac		free
361	P 1	pS G R G K Q G G Kac A R A Kac A K S R S S	H2a 1-19	S1P	K9ac	K13ac		free
362	P 2	S G R G Kac Q G G Kac A R A Kac A K S R S S	H2a 1-19	K5ac	K9ac	K13ac		free
363	P 3	pS G R G Kac Q G G Kac A R A Kac A K S R S S	H2a 1-19	S1P	K5ac	K9ac	K13ac	free
364	P 4	P D P A K S A P A P K K G S K K A V T	H2B 1-19	unmod				free
365	P 5	P D P A Kac S A P A P K K G S K K A V T	H2B 1-19	K5ac				free
366	P 6	P D P A K S A P A P K Kac G S K K A V T	H2B 1-19	K12ac				free
367	P 7	P D P A K S A P A P K K G pS K K A V T	H2B 1-19	S14P				free

368	P 8	P D P A K S A P A P K K G S Kac K A V T	H2B 1-19	K15ac				free
369	P 9	P D P A Kac S A P A P K Kac G S K K A V T	H2B 1-19	K5ac	K12ac			free
370	P10	P D P A Kac S A P A P K K G pS K K A V T	H2B 1-19	K5ac	S14P			free
371	P11	P D P A Kac S A P A P K K G S Kac K A V T	H2B 1-19	K5ac	K15ac			free
372	P12	P D P A K S A P A P K Kac G pS K K A V T	H2B 1-19	K12Ac	S14P			free
373	P13	P D P A K S A P A P K Kac G S Kac K A V T	H2B 1-19	K12Ac	K15Ac			free
374	P14	P D P A K S A P A P K K G pS Kac K A V T	H2B 1-19	S14P	K15Ac			free
375	P15	P D P A Kac S A P A P K Kac G pS K K A V T	H2B 1-19	K5Ac	K12Ac	S14P		free
376	P16	P D P A Kac S A P A P K Kac G S Kac K A V T	H2B 1-19	K5Ac	K12Ac	K15Ac		free
377	P17	P D P A Kac S A P A P K K G pS Kac K A V T	H2B 1-19	K5Ac	S14P	K15Ac		free
378	P18	P D P A K S A P A P K Kac G pS Kac K A V T	H2B 1-19	K12Ac	S14P	K15Ac		free
379	P19	P D P A Kac S A P A P K Kac G pS Kac K A V T	H2B 1-19	K5ac	K12Ac	S14P	K15Ac	free
380	P20	Bio A A N W S H P Q F E K A A	Biotin, control peptide					biotinylated
381	P21	E Q K L I S E E D L A	c-myc tag					free
382	P22	HAc	neg. contol					acetylated
383	P23	K Kme1 Kme2 Kme3 Kac R Rme2s R Rme2a R Cit K Kme1 Kac Kme3 R K	background 01					acetylated
384	P24	R Rme2s K Kme1 Kac R Rme2a Kme2 K Kme3 R Kme1 Rme2s K Kac R K	background 02					acetylated

*CelluSpots™ arrays are manufactured under license by INTAVIS Bioanalytical Instruments AG

Chip 2x duplicate of 384 peptides:

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