In order to gain a more global view of the activity of histone demethylases, we report here genome-wide studies of the fission yeast SWIRM and polyamine oxidase (PAO) domain homologues of mammalian LSD1. Consistent with previous work we find that the two S. pombe proteins, which we name Swm1 and Swm2 (after SWIRM1 and SWIRM2), associate together in a complex. However, we find that this complex specifically demethylates lysine 9 in histone H3 (H3K9) and both up- and down-regulates expression of different groups of genes. Using chromatin-immunoprecipitation, to isolate fragments of chromatin containing either H3K4me2 or H3K9me2, and DNA microarray analysis (ChIP-chip), we have studied genome-wide changes in patterns of histone methylation, and their correlation with gene expression, upon deletion of the swm1\(^{-}\) gene. Using hyper-geometric probability comparisons we uncover genetic links between lysine-specific demethylases, the histone deacetylase Clr6, and the chromatin remodeller Hrp1. The data presented here demonstrate that in fission yeast the SWIRM/PAO domain proteins Swm1 and Swm2 are associated in complexes that can remove methyl groups from lysine 9 methylated histone H3. \textit{In vitro}, we show that bacterially expressed Swm1 also possesses lysine 9 demethylase activity. \textit{In vivo}, loss of Swm1 increases the global levels of both H3K9me2 and H3K4me2. A significant accumulation of H3K4me2 is observed at genes that are up-regulated in the swm1\(^{-}\) strain. In addition, H3K9me2 accumulates at some genes known to be direct Swm1/2 targets that are down-regulated in the swm1\(^{-}\) strain. The \textit{in vivo} data indicate that Swm1 acts in concert with the HDAC Clr6 and the chromatin remodeller Hrp1 to repress gene expression. In addition, our \textit{in vitro} analyses suggest that the H3K9 demethylase activity requires an unidentified post-translational modification to allow it to act. Thus, our results highlight complex interactions between histone demethylase, deacetylase and chromatin remodelling activities in the regulation of gene expression.

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the MS analysis are presented in Table S1 in the Supplementary Information and are summarized in Figure 1. In brief, our data confirm the results of Nicolas et al., (2006) [14] and show that the Swm1 complex contains Swm2 and two new PHD domain containing proteins (S. pombe DB CDS: SPCC4G3.07 and SPAC3D11.08c), hereafter referred to as Swp1 (S. pombe associated PHD1) and Swp2 (S. pombe associated PHD2). Surprisingly, the purified Swm2 complex contained only Swm1 and Swp1, but not Swp2, suggesting that the Swm proteins may exist in more than one complex. However, in contrast to the results of Nicolas et al., (2006) [14] we did not detect either Hrp1 or SPBP738.01 (an RNA recognition motif protein), suggesting that these proteins have weaker affinities or are more transiently associated with the complexes. Alternatively, the use of different tags in the two studies may also explain the discrepancy.

The S. pombe Swm complex demethylates lysine-9 of histone H3

Previous studies have shown that human LSD1 can demethylate either lysine residues 4 or 9 in histone H3, depending on the presence or absence of associated proteins [13]. To determine whether the S. pombe Swm complex can also demethylate histones, we carried out a series of demethylation assays using purified TAP-tagged complexes.

Initial assays for histone demethylase activity were attempted using similar methods to those described by Shi et al., (2004) [2], involving MS and synthetic peptide substrates. However, when comparing the detection limits (assuming a similar level of activity involving MS and synthetic peptide substrates. However, when using similar methods to those described by Shi et al., (2004) [2], we carried out a series of demethylation assays using purified TAP-tagged complexes.

Swp2 (SPAC30D11.08c) are annotated with domain borders as in the SMART sequence analysis database (http://smart.embl-heidelberg.de). doi:10.1371/journal.pone.0000386.g001

Figure 1. Schematic representation of Swm1/2 complex members: Swm1 (SPBC146.09c), Swm2 (SPAC3E2.02), Swp1 (SPCC4G3.07) and Swp2 (SPAC3D11.08c) are annotated with domain borders as in the SMART sequence analysis database (http://smart.embl-heidelberg.de). doi:10.1371/journal.pone.0000386.g001

Interestingly, only methylated H3K9 was found to be a substrate for the S. pombe Swm1 and Swm2 complexes (Figure 2a). Similar levels of activity were detected regardless of whether the complex was purified using TAP-tagged Swm1 or Swm2. Moreover, similar levels of activity were found for histone H3 substrates in the form of either purified bulk histones or nucleosomes. By contrast, in the human LSD1 control we observed specific demethylation of H3K4 as previously reported (Figure 2b) [2]. We next tested whether the S. pombe Swm1 and Swm2 complexes were able to demethylate K9 in isolated histone H3. While both TAP-tagged complexes were found to have similar activity towards histone H3 purified from calf thymus (in comparison to nucleosomal substrates), neither complex was active on recombinant histone H3 (Figure 2c). It is currently unclear why the S. pombe demethylase complex is inactive on recombinant methylated histone H3, but one possibility is that an additional post-translational modification(s) is needed for the complex to effectively recognise its substrate. Finally, to rule out the possibility that a contaminant was responsible for the H3K9 demethylase activity, we demonstrated that recombinant E. coli expressed Smo1 has the same pattern of activity/specificity as that of the intact complex (Figure 2d). This experiment demonstrates that Swm1 is a catalytically active sub-unit. However, because we have not so far been able to express/purify recombinant Swm2, this PAO domain protein may also be catalytically active. Structure-based sequence analysis suggests that Swm2 also has the appropriate FAD-binding and catalytic residues required for demethylase activity (data not shown).

Swm1 both up- and down-regulates gene expression

To determine if the Swm1 and Swm2 proteins are involved in regulation of gene expression, we constructed strains of S. pombe in which either the Swm1 or Swm2 genes were deleted. Consistent with the results obtained by Nicolas et al., (2006) [14], we found that swm2 is an essential gene and that deletion of swm1 markedly increased the cell doubling time (data not shown). Thus, gene expression profiling could only be carried out in cells lacking Swm1.

Our global analysis of gene expression shows that somewhat more genes are up-regulated in the swm1 deletion (swm1A) strain (263 genes) than are down-regulated (173 genes) (using a 1.5 fold cut-off; see Table S2 in the Supplementary information). In Figure 3, the moving average (calculated over 150 genes) of the swm1A/wild-type expression ratio vs the level of wild-type gene expression is plotted. If the swm1 deletion had no effect on gene expression one would expect a global moving average ratio of 1.0. As can be seen, however, the swm1A/wild-type expression ratio is generally high for non-expressed and weakly expressed genes, suggesting that (globally) Swm1 represses these non-expressed and weakly expressed genes.

We emphasise, however, that Swm1 also has an activating role at 173 genes. The effect of swm1 deletion on these genes appears to be direct. In hyper-geometric distribution tests, we found a statistically significant overlap when comparing the published genome-wide Swm1/2 localisation data [14] with the list of genes that are down-regulated in swm1A cells (P = 3.18 × 10^-7), but not those that are up-regulated (compare Figure 4c bottom, left and right).

Further analysis, using hyper-geometric distribution tests, with a database of genes affected by histone modifications, showed that there is a very significant similarity of up-regulated genes in the swm1A strain to up-regulated genes in the chr6-I strain (see Table 1). chr6-I is a loss of function mutant of Clr6, a histone deacetylase
(HDAC) and known transcriptional repressor, which is similar to HDAC1/2 in higher organisms [16–17]. Consistent with Swm1 acting in concert with Clr6, there is also a significant overlap of genes that are up-regulated in the swm1Δ strain with genes whose promoters lack acetylation in wild-type cells [18] (see Table 1).

Our analyses extend the observations of Nicolas et al., (2006) [14]. Thus, it is now clear that the S. pombe Swm1/2 complex has dual roles in gene regulation, up-regulating some 173 genes and down-regulating some 265 others. The correlation between Swm1/2 localisation and down-regulated genes in swm1Δ cells, suggests that the complex might have a more direct role in stimulating gene expression than it does in repression. Moreover, the imposition of a repressed state by Swm1/2 involves one of the HDACs – Clr6.

**The effects of Swm1 deletion on K4 and K9-methylation of histone H3**

We next measured levels of H3K4 and H3K9 dimethylation using ChIP-chip experiments in both inter-genic (IGR) (including all pol II promoters) and open-reading frame (ORF) regions in the swm1Δ strain [see Ref. 18 for details of the microarray platform]. (Our studies focussed on the levels of dimethylation, as opposed to monomethylation, because of the availability of suitable antibodies. In addition, levels of trimethylation are known to be low in S. pombe.) In agreement with our finding that the fission yeast Swm complex demethylates H3K9 (Figure 2), deletion of swm1 resulted in increased levels of H3K9me2 in a large part of the genome – 8.2% of genes showed increased H3K9me2 in either the ORF or IGR regions. H3K4me2 levels, however, were also increased in a smaller part of the genome (3.8%). Interestingly, there appears to be a bias towards increased levels of H3K4me2 in ORF regions (as compared to IGR regions), whereas H3K9me2 levels were increased in both the ORF and IGR regions (compare Figures 4a, 4b). (See Table S3 in the Supplementary information for lists of the IGR and ORF regions with high levels of H3K4me2 and H3K9me2 in the swm1Δ strain.) (Note that the S. pombe genome is composed of roughly equal amounts of IGR and ORF regions.)

Globally, overall levels of H3K9me2 were not significantly changed in genes whose expression is either up- or down-regulated in the absence of Swm1 (Figure 4a). However, on closer examination of the genes that are activated by Swm1, a few have increased H3K9me2 levels in the absence of Swm1. Within this group of 10 genes, stil1+ and SPCC1620.06c were previously identified as Swm1/2 binding targets [14]. This data suggests that in wild-type
cells Swm1 may have an activating function at some genes, which involves demethylation of H3K9me2 in ORF regions, consistent with the general role of H3K9me2 in gene repression and silencing in *S. pombe* [19–21]. However, due to the small number of genes involved, this does not appear to be the major role of the Swm1/2 complex.

In genes that are repressed by Swm1, and therefore up-regulated in the *swm1Δ* strain, levels of H3K4me2 were seen significantly more frequently (CHI square, \( P < 0.01 \)). The bias towards increased levels of H3K4me2 in ORF regions (as compared to the IGR regions) was again apparent (Figure 4b). However, no difference in H3K4me2 levels was observed in genes that are activated by Swm1, and therefore repressed in the *swm1Δ* strain.

The bias towards increased H3K4me2 levels in ORF regions of genes that are up-regulated in *swm1Δ* cells, when taken together with the lack of in-vitro H3K4 demethylase activity of the Swm1/2 complex (see above), suggests that the increased H3K4me2 levels results from either increased H3K4 methylation via Set1, or the incorporation of H3K4 methylated histones during transcription.

**Correlation of H3K9me2 or H3K4me2 levels with other post-translational modifications**

The changes in H3K9me2 or H3K4me2 levels observed at genes in the *swm1A* cells may correlate with other post-translational modifications associated with the same genes. To test this we compared the lists of IGR and ORF regions showing either high H3K9me2 or H3K4me2 in *swm1A* cells with our database of genes affected by histone modifications using hyper-geometric distribution tests.

The regions which show increased H3K4me2 in the *swm1A* cells were very significantly similar to regions where histone acetylation is low in wild-type cells. They were also significantly similar to regions with increased acetylation in the *clr6-1* mutant [18] (Table 2). These findings are consistent with the observation that there is a very significant similarity of up-regulated genes in the *swm1A* strain to up-regulated genes in the *clr6-1* strain (see above and Table 1). The functional link to Clr6 is also interesting given that LSD1 (a human homologue of Swm1) physically and functionally interacts with HDAC1/2 [7–10]. Moreover, the regions associated with increased H3K4me2 in *swm1Δ* cells tend to be the 3’ regions of longer genes (>1000 bp), which in wild type cells normally have low levels of both histone acetylation and H3K4me2, and which are hyperacetylated in *clr6-1* cells [22]. These findings again suggest that Swm1 and Clr6 may collaborate to maintain repressive chromatin in both IGR and ORF regions, to influence both transcriptional initiation and elongation. Furthermore, the affected IGR regions in the *swm1A* cells also overlap significantly with Cse3 and Sir2 HDAC binding data (Table 2).

The lists of IGR regions which show increased levels of H3K4me2 in the *swm1A* strain also overlapped significantly with lists of genes representing IGR binding of the Hrp1 chromatin remodelling factor and its closely related (64% identical) paralogue Hrp3 (\( P = 9.01 \times 10^{-6} \) and \( 5.64 \times 10^{-4} \), respectively; see Table 2). Moreover, the lists of up-regulated genes in the *swm1A* strain showed significant similarity to lists of IGR regions bound by Hrp1 and Hrp3 (\( P = 5.48 \times 10^{-6} \) and \( 1.33 \times 10^{-5} \), respectively; See Table 1).

**Possible targeting of the Swm complex via Clr6?**

Given the finding that loss of function of either the Swm1 histone demethylase or Clr6 HDAC, results in up-regulation of genes and altered levels of histone modifications in the same regions of the
Figure 4. Genome wide analysis of histone methylation and gene expression in swm1 cells, and binding of Swm1/2. A) A comparison of increased H3K9me2 and altered gene expression in swm1 deletion cells. The Venn diagrams illustrate the degree of overlap between lists of IGR and ORF regions having high H3K9me2 in swm1 deletion cells (using a cutoff value of 2.0), and a list of genes, which showed altered expression in swm1Δ (using a cutoff value of 1.5). Left: 8.2% of total S. pombe genes showed high IGR or ORF H3K9me2 levels in swm1Δ. Middle: The Venn diagram shows the fraction of genes up-regulated in swm1Δ having high IGR or ORF H3K9me2. Right: The Venn diagram shows the fraction of genes down-regulated in swm1Δ having high IGR or ORF H3K9me2. The inserted table shows a list of 10 genes down-regulated in swm1Δ, which also show high H3K9me2 ORF levels in swm1Δ (Swm1/2 binding targets are indicated in bold).

B) A comparison of increased H3K4me2 and altered gene expression in swm1 deletion cells. The Venn diagrams illustrate the degree of overlap between lists of IGR and ORF regions having high H3K4me2 in swm1 deletion cells (using a cutoff value of 2.0), and a list of genes, which showed altered expression in swm1Δ (using a cutoff value of 1.5). Left: 3.8% of total S. pombe genes showed high IGR or ORF H3K4me2 levels in swm1Δ. Middle: The Venn diagram shows the fraction of genes up-regulated in swm1Δ having high IGR or ORF H3K4me2 levels, which is significantly more than expected from the genome average (CHI square, P<0.01; indicated). Right: The Venn diagram shows the fraction of genes down-regulated in swm1Δ having high IGR or ORF H3K4me2.

C) Comparison of Swm1/2 binding targets (determined by Nicolas et al., (2006)), H3K4me2 and H3K9me2 levels, as well as gene expression changes in swm1 deletion cells. Top: The Venn diagrams illustrate the degree of overlap between lists of IGR and ORF regions having high H3K4me2 and H3K9me2 levels (as indicated) in swm1 deletion cells, and a list of Swm1/2 binding targets. 13.4% of Swm1/2 binding targets showed high IGR or ORF H3K4me2 levels, which is significantly more than expected from the genome average (CHI square, P<0.001; indicated). Bottom: The Venn diagrams illustrate the degree of overlap between the list of Swm1/2-binding targets and gene expression changes in swm1 deletion cells. Bottom right: A significant proportion of swm1 down-regulated genes (hypergeometric P value indicated) were defined as Swm1/2-binding targets. (Note that our microarray contains only 143 of the 175 targets published by Nicolas et al., (2006)).

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genome, and the knowledge that human LSD1 physically and functionally interacts with HDAC1/2 [7–10], we wondered whether there was a physical interaction between the S. pombe Swm1 and Clr6 complexes. Because we had not identified any of the Clr6 complex components in TAP-tagging/MS studies of Swm1 and Swm2, we also TAP-tagged components of the Clr6 complex, to see if we could identify any of the Swm partner proteins. These studies, however, only identified the same proteins found in previous work [23], namely: Clr6, Alp13, Prw1 and Pst2 (data not shown). In preliminary experiments we have found that Hrp1 and Hrp3 co-purify in affinity purification experiments (Walfridsson, Khorosjutina, Gustafsson, Ekwall et. al. manuscript in preparation). Our results, therefore, indicate that there is a conserved functional interaction between the Swm1 histone demethylase and the Clr6 histone deacytelase that is also found in mammalian cells. They also provide further support for a model in which deacyetation of nucleosomes sets the stage for demethylation [12]. In S. pombe, however, despite the suggested functional association, and unlike the situation in higher organisms, any interaction between the two complexes appears to be of a more transient character.

**Possible targeting of the Swm complex via Hrp1/3?**

Previously, Nicolas et al (2006) [14] showed that the Swm1/2 complex physically interacts with Hrp1. The putative functional links to the Hrp1 and Hrp3 chromatin remodelling factors that we identify here suggest that this interaction may provide a mechanism for targeting of the Swm demethylase. The double chromo

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**Table 1.** Hyper-geometric probability comparisons of genes either down- or up-regulated in *swm1* deletion cells.

| P value | Similar list Name |
|---------|-------------------|
| 1.25×10−33 | stress up >2 fold |
| 3.34×10−29 | HIGH expression in cr6-1 cr3 deletion (1.5 fold) |
| 4.99×10−29 | wild type meiosis up >2 fold |
| 6.30×10−28 | HIGH expression in cr6-1 (1.5 fold) |
| 3.82×10−11 | LOW IGR WT H4K5Ac H3 Cter corr (1.5 fold) |
| 8.42×10−09 | LOW IGR WT H4K12Ac H3 Cter corr (1.5 fold) |
| 1.00×10−06 | LOW IGR WT H3K9Ac H3 Cter corr (1.5 fold) |
| 5.48×10−06 | IGR binding of Hrp1 (0.90 percentile) |
| 8.92×10−06 | HIGH expression in cr3 deletion (1.5 fold) |
| 1.33×10−05 | IGR binding of Hrp3 (0.94 percentile) |
| 7.14×10−05 | LOW IGR WT H4K16Ac H3 Cter corr (1.5 fold) |
| 0.0000185 | LOW IGR WT H4K14Ac H3 Cter corr (1.5 fold) |
| 0.000022 | Process-Carbohydrate Metabolism |
| 0.0196 | ORF binding Clr3-myc (0.86 percentile) |

**LOW Expression in *swm1* vs wild-type, cut-off 1.5 (3 of 4), 173 genes**

| P value | Similar list Name |
|---------|-------------------|
| 2.03×10−07 | HIGH expression in wild type |
| 0.00014 | Process-Amino Acid Metabolism |
| 0.00418 | GO: 0006531: Process: aspartate metabolism |
| 0.0101 | GO: 0006555: Process: methionine metabolism |
| 0.0117 | Process-Sulfur Metabolism |
| 0.0496 | Process-Transporters |

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**Table 2.** Hyper-geometric probability comparisons of IGR and ORF lists with high H3K4me2 in *swm1* deletion cells.

| P value | Similar list Name |
|---------|-------------------|
| 1.64×10−19 | High IGR clr6-1 H4K5Ac H3 cter corr (2 fold) |
| 2.33×10−18 | LOW IGR WT H3K14Ac H3 Cter corr (1.5 fold) |
| 2.82×10−16 | High IGR clr6-1 H4K12Ac H3 cter corr (2 fold) |
| 1.82×10−15 | High IGR clr6-1 H3K14Ac H3 cter corr (2 fold) |
| 5.59×10−15 | LOW IGR WT H4K5Ac H3 Cter corr (1.5 fold) |
| 7.23×10−15 | LOW IGR WT H3K9Ac H3 Cter corr (1.5 fold) |
| 7.31×10−13 | LOW IGR WT H4K12Ac H3 Cter corr (1.5 fold) |
| 7.64×10−13 | HIGH IGR clr6-1 H4K16Ac H3 cter corr (2 fold) |
| 6.45×10−12 | LOW IGR WT H3K4me2 H3 cter corr (1.5 fold) |
| 2.11×10−09 | LOW IGR WT H4K16Ac H3 Cter corr (1.5 fold) |
| 6.05×10−08 | High IGR clr6-1 H3K9Ac H3 cter corr (2 fold) |
| 9.01×10−06 | Hrp1 IGR binding (0.9 percentile) |
| 0.000564 | Hrp3 IGR binding (0.94 percentile) |
| 0.00058 | HIGH expression in wild type meiosis (2 fold) |
| 0.00303 | IGR Binding Sir2-myc (0.91 percentile) |
| 0.0497 | IGR binding Clr3-myc (0.86 percentile) |

**HIGH ORF H3K4me2 in *swm1*, 162 genes**

| P value | Similar list Name |
|---------|-------------------|
| 2.65×10−28 | LOW IGR WT H3K14Ac H3 Cter corr (1.5 fold) |
| 2.4×10−23 | LOW IGR WT H4K5Ac H3 Cter corr (1.5 fold) |
| 3.07×10−20 | LOW IGR WT H4K16Ac H3 Cter corr (1.5 fold) |
| 3.65×10−20 | Gene length (>1000) |
| 6.0×10−18 | LOW IGR WT H4K12Ac H3 Cter corr (1.5 fold) |
| 1.1×10−09 | Gene length (>2000) |
| 4.31×10−06 | Hrp3 ORF binding (0.92 percentile) |
| 6.66×10−06 | LOW IGR H3K4me2 H3 Cter corr (1.5 fold) |
| 2.99×10−05 | LOW IGR WT H4K12Ac H3 Cter corr (1.5 fold) |
| 5.46×10−05 | LOW IGR WT H3K9Ac H3 Cter corr (1.5 fold) |
| 0.000123 | Swm1/2 binding targets |
| 0.000535 | High IGR clr6-1 H4K12Ac H3 cter corr (2 fold) |
| 0.0629 | High IGR clr6-1 H4K16Ac H3 cter corr (2 fold) |
| 0.0324 | High IGR clr6-1 H4K5Ac H3 cter corr (2 fold) |
| 0.0455 | High IGR clr6-1 H3K14Ac H3 cter corr (2 fold) |

**HIGH IGR or ORF H3K4me2 in *swm1*, combined list 199 genes**

| P value | Similar list Name |
|---------|-------------------|
| 0.0032 | Swm1/2 binding targets |
| 0.0133 | HIGH Expression in *swm1* (1.5 fold) |

do可观测到，Swm1和Clr6 complexes的相互作用是通过Hrp1/3实现的。然而，这种相互作用在更高的生物体中并未观察到。因为Hrp1和Hrp3共纯化，我们推测Hrp1/Swm complexes的相互作用可以通过与Hrp3的相互作用来实现。
Concluding remarks
In conclusion, the data presented here demonstrate that in fission yeast the SWIRM/PAO domain proteins Swm1 and Swm2 are associated in complexes that can remove methyl groups from lysine 9 methylated histone H3. Interestingly, recognition of the H3K9 substrate appears to require an as yet unidentified modification. In vivo, loss of Swm1 increases the global levels of H3K9me2 and H3K4me2, and it results in a significant accumulation of H3K4me2 at genes that are up-regulated in the swm1Δ strain. The bias towards increased H3K4me2 levels in ORF regions of genes that are up-regulated (as compared to IGR regions), suggests that this increase results from either increased methylation by Set1, or incorporation of H3K4 methylated histones, during transcription.

An alternative explanation for the increased levels of H3K4me2 is that Hrp1/3 might act as coregulators, and influence the specificity of the Swm1 complex. In a similar manner to that of human LSD1, which when complexed with the androgen receptor switches from a H3K4 to a H3K9 demethylase [13], it is possible that K4 demethylation is favored over K9 in the functional context of the chromatin remodelling factors Hrpl and Hrp3. However, the bias to increased levels of H3K4me2 in ORF regions, suggests (as discussed above) that the increased levels at these genes results in some way from increased transcription.

Aside from at a few genes, where in the swm1Δ deletion increased H3K9me2 levels are correlated with reduced levels of gene expression, the functional role of the Swm1/2 H3K9 demethylase activity is not yet clear. Our results, however, highlight complex interactions between histone demethylase, histone deacetylase and chromatin remodelling activities in the regulation of gene expression. The in vivo data indicate that Swm1 acts in concert with the HDAC Clr6 and the chromatin remodeller Hrp1 to repress gene expression, but further work is necessary to uncover the nature of these functional interactions.

Supplementary information is available on line.

METHODS
Reagents
IgG Sepharose was from Amersham, Ni-magnabeads were from Promega and the Dynabeads M280 were from Dynal Biotech. Rabbit anti-mouse immunoglobulin protein used for coating the beads was from DAKO, A/S Denmark. Calf thymus bulk histones and histone H3 were purchased from Roche. Chicken poly-nucleosomes were purchased from Abcam and recombinant Xenopus laevis H3 was from Upstate.

Construction of strains
Strains expressing TAP-tagged Swm1 and Swm2, and the swm1Δ knockout strain, were constructed using previously described protocols [26–28]. In brief, the TAP-tagging constructs for homologous recombination were made using fusion PCR of 500 base pair genomic DNA fragments and the C-terminal TAP-tagging cassette. The resulting DNA fragment was transfected into S. pombe strain 501 (leu1-32; ura4-D18; ade6-704/h−). Homologous recombination was confirmed by PCR with primers inside and outside the incorporated TAP-tag. Expression of the construct and fusion protein solubility was checked by Western blots with anti-protein A antibodies. Strain viability was compared to that of the wild type strain in normal growth and under stress conditions (37°C). The swm1Δ knockout strain was constructed by homologous integration of Clonat, replacing the ORF. The integration was confirmed by PCR using primers 5’ and 3’ to the recombined region together with primers internal to the Clonat gene.

Cells growth and affinity purification of the complexes
The published TAP-purification procedure [29] was followed with minor changes. The wild type and TAP-tagged strain cells were harvested and washed in ice-cold water, resuspended in 20 mM HEPES pH 8.0, 150 mM NaCl, 0.1% Tween 20 and broken in liquid nitrogen using a SPEX CerilliPrep 6500 Freezer Mill. The soluble fraction was bound to either IgG coated Sepharose or IgG coated Dynabeads, washed extensively, cut with His-tagged TEV protease and eluted from the beads. Excess IgG and TEV were removed by incubation with protein A and Ni-agarose beads. The eluate was bound to calmodulin-binding beads and eluted with EGTA containing buffer. To avoid loss of protein the complex was not further purified by SDS-PAGE. Instead, the eluted complex was directly digested with trypsin and the resulting peptides identified by mass-spectrometry.

Purification of hLSD1
Human LSD1 (residues 72-852) was subcloned into the pET30 vector (Novagen). Protein expression was carried out in the E. coli Rosetta 2 strain (Novagen). To induce expression 0.5 mM IPTG was added to cultures at OD600 of 0.6 and incubated at 25°C for 4 hours. Harvested cells were disrupted in lysis buffer (40 mM Tris pH 8.0, NaCl 300 mM, 0.5% NP40, protease inhibitors (Sigma)) using an Emulsiflex-05 (Avestin) at up to 8000 psi. hLSD1 was purified from the clarified lysate with Ni-NTA agarose (Qiagen) and further purified by anion exchange chromatography (MonoQ column (Pharmacia Biotech)) and gel filtration (Superdex 200 (Pharmacia Biotech)) using standard protocols.

Purification of GST-Swm1
Full length GST-Swm1 was subcloned into pGEX-2T. Protein expression was carried out in the E.coli Rosetta strain. To induce expression 0.5 mM IPTG was added to cultures at an OD600 of 0.5 and incubated at 15°C for 16 hours. Harvested cells were disrupted in lysis buffer (25 mM Tris-HCl pH 8.0, 150 mM NaCl, 0.1% Triton-X-100, protease inhibitors (Roche)) by sonication. Glutathione sepharose resin (Pharmacia) was added to the clarified lysate and incubated with rotation for 2 hrs at 4°C. The resin was then extensively washed with lysis buffer without Triton-X-100. Bound proteins were finally eluted in 50 mM Tris pH 8.0, 10 mM glutathione.

In-vitro demethylase assays
3H-labelled methyl histone substrates were prepared using the following histone methyltransferases (HMT): GST-Set7, MBP-Clr4 and GST-Set2 as previously described [15]. Briefly, 100 ul labelling reactions were carried out in methylate buffer (50 mM Tris pH 8.0, 1 mM DTT, 10% glycerol and 10 mM ZnCl2) containing 1-5 ug HMT’s, 5 µl S-adenosyl-[3H]methyl-methionine (70 Ci/mmol, NEN) and substrate (50 µg calf thymus bulk histones, 10 µg chicken poly-nucleosomes, 10 µg calf thymus histone H3 or 10 µg recombinant Xenopus laevis H3). After a 5 minute incubation at 37°C, reactions were immediately dialysed into demethylase buffer (50 mM Tris pH 8.8, 0.5 mM DTT and 5% glycerol) at 4°C.

For demethylase assays labelled histone substrates containing 5,000-50,000 cpm were incubated with 25 ul of either TAP-tagged Swm complex, mock purifications from wild-type cells, or 5 µg hLSD1 or GST-Swm1 in a final volume of 100 ul with demethylase buffer at 37°C for 1 hr. The Nash method was then used to detect the formation of 3H-labelled formaldehyde [30].
After precipitation with 10% TCA an equal volume of Nash reagent (3.99 M ammonium acetate, 0.1 M acetic acid and 0.2% 2,4-pentanedione) was added to the supernatant and incubated at 37°C for 50 minutes, followed by extraction with an equal volume of 1-pentanol. The extracted supernatant was then subjected to scintillation counting.

**Genome wide analysis**

The microarray analysis in this study was performed essentially as outlined in [18]. cDNA expression profiling was carried out according to Xue et al., (2004) [31]. We used S. pombe ORF and combined IGR+ORF spotted microarrays from Eurogentec custom DNA microarray services (Belgium). For histone methylation maps, ChiP-chip experiments were carried according to Robyr and Grunstein (2003) [32]. Antibodies recognising specific methylation marks were employed: H3K9me2 and H3K4me2 (a kind gift from Prof. David Allis). For Hrp1 and Hrp3 binding studies we used the ChiP-chip procedure described by Kurdistani et al. (2002) [33]. For expression profiling and analysis of histone methylation levels, two microarray experiments were performed using two independent ChiP samples and including Cy3/Cy5 dye swaps. Because each Eurogentec microarray yields two data points we thus measured four data points for each experiment. Expression profiling and histone methylation in mutant as, wild-type data sets were normalized using the GeneSpring software and Lowess (per spot, per chip) intensity-dependent normalization, which corrects for nonlinear rates of dye incorporation. Cut off values of 1.5 (for gene expression) and 2.0 (for histone methylation levels) in at least 3 out of 4 data points were used to generate the ‘high’ and ‘low’ gene lists. For Hrp1 and Hrp3-binding experiments three microarrays were used, yielding six data points. Here we first used the GeneSpring software for a ‘per chip’ normalization and then employed a statistically determined cut-off for binding according to the median percentile ranking method [34]. Similar gene lists were identified using the automatic hyper-geometric distribution tests in the Gene List inspector function of GeneSpring. The hyper-geometric distribution test calculates the probability of overlap corresponding to k or more IGR or ORF fragments between an IGR or ORF list of m fragments compared against another gene list of n fragments when randomly sampled from a universe of u genes:

$$\frac{1}{u} \binom{m}{i} \binom{n-i}{m-i}$$

The significantly overlapping gene lists were illustrated using Venn diagrams.

**SUPPORTING INFORMATION**

**Table S1** a) Identification of peptides in the Swm complex purified using TAP-tagged Swm1 (SPBC146.09C) b) Identification of peptides in the Swm complex purified using TAP-tagged Swm2 (SPAC23E2.02)

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**Table S2** A list of genes either down- or up-regulated in swm1 deletion cells

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**Table S3** Genes that have either increased levels of H3K4me2 or H3K9me2 in swm1 deletion cells

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The microarray data has been deposited in the GEO database (accession number GSE6224).

**Author Contributions**

Conceived and designed the experiments: RA EL KE DL. Performed the experiments: KE MO DL CB. Analyzed the data: EL KE MO DL CB. Wrote the paper: RA EL KE DL. Other: Contributed to the experimental work and analysis of data: TK JW ST AB JC NM. Contributed to the writing of the paper: NM. Provided assistance with the TAP-tagging experiments: AC PW.

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