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Combinatorial readout of unmodified H3R2 and acetylated H3K14 by the tandem PHD finger of MOZ reveals a regulatory mechanism for HOXA9 transcription

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Histone acetylation is a hallmark for gene transcription. As a histone acetyltransferase, MOZ (monocytic leukemia zinc finger protein) is important for HOX gene expression as well as embryo and postnatal development. In vivo, MOZ forms a tetrameric complex with other subunits, including several chromatin-binding modules with regulatory functions. Here we report the solution structure of the tandem PHD (plant homeodomain) finger (PHD12) of human MOZ in a free state and the 1.47 Å crystal structure in complex with H3K14ac peptide, which reveals the structural basis for the recognition of unmodified R2 and acetylated K14 on histone H3. Moreover, the results of chromatin immunoprecipitation (ChIP) and RT–PCR assays indicate that PHD12 facilitates the localization of MOZ onto the promoter locus of the HOXA9 gene, thereby promoting the H3 acetylation around the promoter region and further up-regulating the HOXA9 mRNA level. Taken together, our findings suggest that the combinatorial readout of the H3R2/K14ac by PHD12 might represent an important epigenetic regulatory mechanism that governs transcription and also provide a clue of cross-talk between the MOZ complex and histone H3 modifications.

[Keywords: histone acetyltransferase; MOZ; PHD finger; H3R2me0; H3K14ac; HOXA9]

Supplemental material is available for this article.

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The eukaryotic genome is assembled into chromatin. The basic unit of chromatin is the nucleosome, which is comprised of 147 base pairs [bp] of DNA wrapped around the histone octamer [two copies each of histones H2A, H2B, H3, and H4] (Kornberg and Lorch 1999). Post-translational modifications [PTMs] such as acetylation and methylation of histones play important regulatory roles in almost every aspect of DNA metabolism, including transcription, replication, recombination, and DNA repair (Berger 2007; Downs et al. 2007; Groth et al. 2007; Kouzarides 2007). Among all of the modifications, histone acetylation at the promoter regions is a hallmark for an actively transcribed gene, and the level of acetylation is positively correlated with transcription rates (Liu et al. 2005; Pokholok et al. 2005; Rosaleny et al. 2007; Wang et al. 2008). The general mechanisms could be categorized into cis and trans. In a cis mechanism, histone acetylation is believed to relax chromatin via the neutralization of positive charges on lysines, thereby reducing the interaction between histones and DNA. On the other hand, a trans mechanism relies on the recruitment of epigenetic readers with various biological functions, as long as other effector proteins associated with them (Kouzarides 2007; Wang et al. 2007).

Monocytic leukemia zinc finger protein [MOZ, also referred to as KAT6A and MYST3] is a MYST family histone acetyltransferase [HAT] that catalyzes the transfer of an acetyl group from acetyl-CoA to the lysine ε-amino groups on histones, which plays roles in hematopoiesis (Thomas et al. 2006), skeletogenesis (Miller et al. 2004; Crump et al. 2006), and other developmental processes.
It is a coactivator of various transcription factors [particularly for hematopoietic specificity] such as RUNX1 (AML1 [acute myeloid leukemia 1]) and PU.1 (Kitabayashi et al. 2001; Pelletier et al. 2002). MOZ was first identified in AMLs through its presence in reciprocal chromosomal translocations, leading to the fusion of MOZ to a transcription activator, CBP [Borrow et al. 1996]; CBP-like coactivator p300 (Chaffanet et al. 2000); or a nuclear receptor coactivator, TIF2 (Carapeti et al. 1998). Each of these fusion proteins contains one or more HAT domains, indicating a role of aberrant acetylation in oncogenic transformation (Yang and Ullah 2007). Moreover, MOZ has been reported to be indispensable in the maintenance of hematopoietic stem cells and responsible for the differentiation of erythroid and myeloid cells [Katsumoto et al. 2006]. Recently, another study in mouse embryos shows that MOZ is required for normal levels of H3K9 acetylation and gene expression at a large number of Hox loci and for identity specification of 19 body segments of the cervical and thoracic axial skeleton and nervous system (Voss et al. 2009).

In vivo, MOZ forms a tetrameric complex with ING5 [inhibitor of growth 5], EAF6 [Esa1-associated factor 6 ortholog], and the bromodomph PID [plant homeodomain] finger proteins BRPF1, BRPF2, or BRPF3 to perform its functions [Ullah et al. 2008]. As the catalytic subunit of the quartet complex, the biological consequences of MOZ depend on its localization to specific gene loci. Insight into how MOZ is recruited to these regions is important for disclosing the mechanisms underlying MOZ-mediated processes. MOZ is a multidomain protein containing a NEMM [N-terminal part of Enok, MOZ, or MORF] domain, a tandem PHD zinc finger, a MYST [HAT] domain for acetyl-transfer catalysis, a glutamate/aspartate (ED)-rich region, and a serine/methionine (SM)-rich region for transcriptional activation [Fig. 1A; Kitabayashi et al. 2001]. The PHD zinc fingers are the only putative histone recognition motif capable of “reading” various histone modifications and unmodified histone tails (residues 1–18) (Fig. 1B,C), but not H4 with Lys16 acetylation (residues 1–18) [Fig. 1B,C], but not H4 with Lys16 acetylated [residues 11–21] [Supplemental Fig. S1], which is slightly different from that of DPF3b. Shortening the H3 tail [residues 1–8] or removing acetylation on H3K14 reduced binding affinity compared with the H3K14ac peptide [residues 1–18] [Fig. 1C; Supplemental Fig. S2], indicating that acetylation of Lys14 could strengthen the interactions between histone H3 and PHD12. Moreover, trimethylation of H3K4 impedes the binding, while acetylation or trimethylation of H3K9 did not show a significant effect [Fig. 1C; Supplemental Fig. S3]. In contrast, methylation of H3R2 devastates the binding completely [Fig. 1D], indicating the essentiality of H3R2 in this binding process. Taken together, these findings suggest that among various histone modifications, the tandem PHD finger of MOZ preferentially recognizes unmodified R2 and acetylated K14 on histone H3.

Considering this bispecificity as independent events, we were curious to study them one by one to figure out how much each site contributes to the overall recognition specificity. Interestingly, although H3 [1–18] is a stronger substrate than H3 [1–8] [Fig. 1C; Supplemental Fig. S2], H3 [9–18] alone does not suffice to induce detectable binding, which could not be rescued by Lys14 acetylation on this fragment [H3 [9–18]K14ac] [Fig. 1C; Supplemental Fig. S6]. Likewise, the destructive effect of alanine sub-

MOZ PHD12 is a H3R2- and H3K14ac-binding module

Results

PHD12 recognizes unmodified R2 and acetylated K14 on histone H3

Bromodomain has long been the only protein domain ever reported to recognize acetyl-lysine residues on proteins, such as GCN5 bromodomain recognizing acetylated Lys16 on histone H4 [Owen et al. 2000], and P/CAB bromodomain binding to lysine-acetylated peptides derived from major acetylation sites on histones H3 or H4 [Dhalluin et al. 1999], until recent studies reported the binding of the tandem PHD finger in D4 protein family member 3b [DPF3b] to acetylated histones [Lange et al. 2008; Zeng et al. 2010]. As the sequence of PHD12 in MOZ is highly similar to that of DPF3b, it is conceivable that MOZ PHD12 is also a candidate for acetylated histone recognition. To address this issue and beyond, we examined the binding affinity of MOZ PHD12 to a collection of histone peptides with or without modifications by NMR [nuclear magnetic resonance] and ITC [isothermal titration calorimetry]. The results showed that MOZ PHD12 binds specifically to histone H3 with Lys14 acetylated [residues 1–18] [Fig. 1B,C], but not H4 with Lys16 acetylated [residues 11–21] [Supplemental Fig. S1], which is slightly different from that of DPF3b. Shortening the H3 tail [residues 1–8] or removing acetylation on H3K14 reduced binding affinity compared with the H3K14ac peptide [residues 1–18] [Fig. 1C; Supplemental Fig. S2], indicating that acetylation of Lys14 could strengthen the interactions between histone H3 and PHD12. Moreover, trimethylation of H3K4 impedes the binding, while acetylation or trimethylation of H3K9 did not show a significant effect [Fig. 1C; Supplemental Fig. S3]. In contrast, methylation of H3R2 devastates the binding completely [Fig. 1D], indicating the essentiality of H3R2 in this binding process. Taken together, these findings suggest that among various histone modifications, the tandem PHD finger of MOZ preferentially recognizes unmodified R2 and acetylated K14 on histone H3 tails.

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stitution of R2 is also not able to be reversed by a single acetylation on Lys14 [H3 (1–18)K14ac_R2A] (Fig. 1E). Therefore, in all respects, only when the hydrogen bonds responsible for R2 recognition remain intact could PHD12 show a preference for acetylated K14 to an unmodified H3 peptide.

Figure 1. PHD12 recognizes unmodified R2 and acetylated K14 on histone H3. (A) Schematic representation of the domain structure of human MOZ. Domains are labeled as follows: (NEMM) N-terminal part of Enok, MOZ, or MORF; (PHD) plant homeodomain-linked zinc finger; (ED) glutamate/aspartate-rich region; (SM) serine/methionine-rich domain. (B) Seven superimposed 1H-15N HSQC spectra of PHD12 [0.3 mM], collected during titration of the H3K14ac peptide, are color-coded according to the molar ratio of peptide PHD12. The red dashed circles indicate the residues with intermediate or slow exchange properties. (C) Dissociation constants of the interactions between PHD12 and histone H3 tail peptides determined by ITC experiments. (D) Superimposed 1H-15N HSQC spectra of PHD12 collected during titration of unH3 [1–18]. (E, top) Superimposed 1H-15N HSQC spectra of PHD12 collected during titration of H3 [1–18]R2me2s or H3 [1–18]R2me2a peptide from left to right, respectively. (Bottom) Chemical shift perturbations in the two-dimensional 1H-15N HSQC spectra and ITC curves introduced by the titration of H3 [1–18]K14ac_R2A.
Overall structures of PHD12 in the free and H3K14ac-bound states

To unveil the mechanism of this combinatorial readout, we solved the solution structure of MOZ PHD12 in a free state (Protein Data Bank [PDB] ID 2LN0) (Fig. 2A) and the crystal structure of that in complex with a histone H3 [1–18] peptide containing R2me0 and K14ac (PDB ID 3V43) (Fig. 2B–D) with structural statistics listed in Supplemental Tables S1 and S2, respectively. The structures showed that each PHD finger adopts a canonical PHD finger fold, which consists of a two-strand anti-parallel β sheet followed by a C-terminal α helix and two zinc atoms coordinated by the Cys4-His-Cys3 motif (Fig. 2B). In each structure, two PHD fingers associate with one another intimately in a face-to-back manner through the interactions between Glu247 and Arg251 in α1 from the first PHD finger (PHD1) and through the interactions between Ser283 and Arg286 in the β3–β4 sheet from the second PHD finger (PHD2). In detail, via the carboxyl and carbonyl oxygen of Glu247, PHD1 forms two hydrogen bonds with two water molecules, bridging the polar contact to PHD2 through the hydroxyl and carbonyl oxygen of Ser283. Similar to the oxygen-mediated hydrogen bond network, a nitrogen atom on the side chain of Arg251 forms a pair of hydrogen bonds with another two water molecules, connecting the two amino nitrogen atoms on the Arg286 side chain (Fig. 2C). These polar interactions immobilize these two PHD fingers to a relatively stationary orientation, making them an integrated structural unit.

PHD2 specifically recognizes unmodified R2 on histone H3

Further analysis of the 1.47 Å resolution complex structure revealed the structural basis for site-specific recognition of unmodified R2 on histone H3 by PHD2. As shown in Figure 2D, the first four residues of the bound H3 peptide (H3K14ac) form a β sheet with α1 of PHD1, and the β3–β4 sheet of PHD2. This interaction results in the formation of additional hydrogen bonds between the amino acids on the peptide and the side chains on PHD2, stabilizing the complex.

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**Table 1:**

<table>
<thead>
<tr>
<th>MOZ PHD12</th>
<th>Kd value (μM)</th>
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<tr>
<td>WT</td>
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<tr>
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</tr>
<tr>
<td>N235A</td>
<td>184.2 ± 64.6</td>
</tr>
<tr>
<td>S210D+N235R</td>
<td>21.4 ± 1.2*</td>
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<td>E261A</td>
<td>65.8 ± 23.1</td>
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<tr>
<td>A275D</td>
<td>5.2 ± 0.8</td>
</tr>
<tr>
<td>D282A</td>
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</tr>
</tbody>
</table>

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**Figure 2:** Structures of PHD12 in the free and H3K14ac-bound states. (A) Ensemble of the 20 lowest-energy structures for residues 204–313 of PHD12; the zinc atoms are highlighted as purple spheres. (B) Cartoon representation of the 1.47 Å crystal structure of PHD12 in H3K14ac-bound states. PHD1 and PHD2 are colored in cyan and blue, respectively. The histone H3 peptide (yellow) and the bound acetate (green) are shown as sticks. (C) PHD1 and PHD2 associate with each other through a series of hydrogen bonds between α1 and the β3–β4 sheet. Intermolecular interactions are depicted as magenta-colored dashed lines. A bridging water molecule involved in intermolecular recognition is shown as a red sphere. (D,E) PHD12 recognition of the N-terminal residues of the H3K14ac peptide, depicted in a protein electrostatic potential surface (D) and ribbon representation (E). The bound H3 peptide and acetate are colored in yellow and green, respectively, with interacting residues on PHD12 colored in blue. (F) Cartoon representation of the superposition of the structures of PHD12 in the free (silver) and H3K14ac peptide-bound (blue) states. The H3 [1–5] segment of the bound peptide is shown in yellow. (G) Dissociation constants of the interaction between PHD12 or mutants and H3K14ac peptides determined by FPA (fluorescence polarization assay) using a C-terminal fluorescein isothiocyanate (FITC)-labeled H3K14ac peptide [residues 1–18] or ITC (indicated with asterisks). (H) In vitro peptide pull-down assays using various biotinylated histone peptides containing the indicated modifications. GST-PHD12 or the mutants were incubated with biotinylated histone peptides immobilized onto streptavidin Sepharose beads. Bound proteins were subjected to SDS-PAGE and stained by Coomassie blue.
H3 peptide are embedded extendedly into the acid groove on the surface of PHD2. More precisely, the N terminus of the peptide is tethered to the main carbonyl oxygen of Gly303, and the side chain of Ala1 in the H3 peptide is buried within a hydrophobic cage formed by residues Met300, Pro301, and Trp305. Located near a patch of negatively charged area, the side chain of Arg2 forms extensive hydrogen bonds with the oxygen atoms on the side chain of Asp282 and Asp285, both of which reside in the hairpin linking β1 and β2 of PHD2. In addition, the Nα2H proton of Arg2 also forms a polar contact with the backbone carbonyl oxygen of Cys281. It is the network of these five hydrogen bonds that stringently guarantees the specific recognition of the H3K2 mark by PHD2. From the structure, we speculated that methylation of R2 will impede the formation of the hydrogen bond network, thus disabling the interaction. Expectedly, the binding of PHD12 to H3 peptide was totally abolished when Arg2 is either symmetrically or asymmetrically dimethylated (Fig. 1E). Besides these, many other polar contacts were observed between PHD2 and the bound H3 peptide: The hydroxyl oxygen of Thr3 interacts with the acylamide group of Asn277; the backbone nitrogens of Lys4 interacts with the carbonyl oxygen of Met278; the Nα nitrogen of Lys4 forms hydrogen bonds with three oxygen atoms from Glu261 and Asn274, mediated by an ordered water molecule; and the acylamide nitrogen of Gln5 interacts with two carbonyl oxygens from the main chain of Asn274 and Asp276, respectively (Fig. 2E). These interactions, together with the hydrogen bond network between PHD2 and the N terminus of the H3 peptide, demonstrate at atomic resolution how recognition of R2 in a H3 sequence context is guaranteed by PHD2.

To further confirm the role of R2 in H3 binding, we used an H3 [1–18]K14ac peptide containing a R2A mutation to titrate PHD12 and monitored it by ITC and NMR, respectively. The results showed no apparent binding when Arg2 was mutated to Ala (Fig. 1E), recapitulating the destructive effect of R2 methylation. Accordingly, several point mutations on PHD2 were prepared: E261A or D282A was designed to disrupt the hydrogen bond formation with Lys4 or Arg2, respectively, while A275D was expected to form an extra polar contact between the residue 275 and Gln5 (Fig. 2E). The FPA (fluorescence polarization assay) results showed that the E261A mutation decreased the binding affinity to the H3K14ac peptide about fourfold to fivefold, and the D282A mutation totally abolished binding, while, on the other hand, the A275D mutant strengthened this binding with a dissociation constant decrease of threefold to fourfold (Fig. 2G). A similar result was obtained from a peptide pull-down assay (Fig. 2H), which further supports the proposed structural basis for H3R2/K4 recognition by PHD2.

From the structure comparison of MOZ PHD12 in the free and H3-bound states (Fig. 2F), the backbones of the two structures are superimposed strictly with a root-mean-square deviation (R.m.s.d.) of 0.77 Å, whereas a conformational change was observed on the segment from Asp269 to Asn277 upon H3 peptide binding. In the free state, this segment folds into a short loop followed by a small α helix, while in the peptide-bound state, it is stretched to form a longer loop to accommodate the side chains of Lys4 and Gln5 (Fig. 2F). The NMR relaxation parameters showed that the segment between Gly272 and Asn277 is relatively flexible (Supplemental Fig. S5), suggesting an “induced fit” of PHD2 during the peptide-binding process.

PHD1 is a histone H3 acetylated Lys14-binding module

Unexpectedly, in the complex structure, instead of acetylated Lys14, an acetate group from the solvent was observed at the acetyl-binding site in PHD1 (Fig. 3C). Furthermore, a side chain of arginine from another PHD12 molecule in the same unit cell occupied the space above the acetate group (Supplemental Fig. S4). Therefore, the competition of acetate in solvent and crystal packing force made us fail to obtain the electron density map of the acetylated Lys14 in the H3K14ac-bound state. In order to investigate the structural basis for acetylated Lys14 recognition by PHD12, a NMR chemical shift perturbation experiment was employed with two histone H3 tail peptides in different Lys14 acetylation states. The comparison between chemical shift changes induced by the addition of different H3 peptides helped us map the acetylated Lys14-binding site on MOZ PHD1. The results summarized in Figure 3A indicated that the overall chemical shift perturbation introduced by H3 [1–18]K14ac is larger than that introduced by H3 [1–18], and the ITC data shown in Figure 3B are consistent with this. Several residues around Ser210 and Asn235 exhibited large chemical shift changes when titrated by H3 [1–18]K14ac (enclosed by red circles in Fig. 3A,C) without being affected by unmodified H3 [1–18], indicating that Ser210 and Asn235 play specific roles in the acetylated Lys14 recognition. Taken together, MOZ PHD1 recognizes acetylated Lys14 on H3 through the collaborative contributions of Ser210, Asn235, and the other residues around them.

In a recent work, Zeng et al. (2010) reported the molecular mechanism underlying the sequence-specific recognition of H3K14ac by DPF3b tandem PHD finger as a functional unit. A DALI (Holm and Rosenstrom 2010) search using MOZ PHD12 as query structure indicated that the tandem PHD finger of DPF3b is the one with the highest structural similarity in all PDB entries, with a Cα r.m.s.d. of 1.9 Å and a Z-score of 13.0. Based on the structural similarity and the same binding substrate, a structural comparison between the tandem PHD fingers of MOZ and DPF3b (Fig. 3D) would be very informative to determine the acetylated Lys14-binding sites on MOZ, which are otherwise invisible in our complex structure. As is demonstrated in the complex structure of DPF3b with the H3K14ac peptide, the methyl group of the acetylated Lys14 inserts into a hydrophobic pocket formed by residues Phe264, Leu296, Trp311, and Ile314. The corresponding hydrophobic residues on MOZ are Phe211, Leu242, Trp257, and Ile260, which together indeed make up a conserved hydrophobic pocket. Moreover, in the DPF3b complex, the rim of the binding pocket is clustered with Asp263, Arg289, and zinc-coordinating Cys313, with the
Figure 3. MOZ PHD1 is a histone H3 acetylated Lys14-binding module. (A) The histogram displays $^1$H, $^15$N chemical shift changes caused by H3 [1–18] (red) or H3 [1–18]K14ac (green) peptide binding. Residues S210, N235, and K243 are circled by red dashed lines. (B) Superposed exothermic ITC enthalpy plots for the binding of PHD1 to H3 [1–18] and H3 [1–18]K14ac peptides. The insert lists the measured binding constants. (C) The molecular surface of PHD1 with residues colored according to the magnitude of the observed chemical shift change upon addition of H3K14ac peptide. The surface residues are color-coded according to the following scheme: (blue) $\Delta d > 0.6$; (marine) $0.6 > \Delta d > 0.4$; (light purple) $0.4 > \Delta d > 0.2$. The positions of residues S210, N235, K243, D276, L279, and D282 are shown. The bound peptide (yellow) and acetate group (green) are shown in sticks. The red dashed circle in C encloses the residues in the first PHD finger that are involved in acetylated lysine recognition. (D, left) Superimposition of the structures of tandem PHD fingers of MOZ (blue) and DPF3b (salmon) in complex with H3K14ac peptides (yellow in the MOZ complex and orange in the DPF3b complex). (D, right) Chemical shift perturbation of residue Lys243 showing that Lys243 takes a role in the interaction with T11 on histone H3K14ac peptide. (E,F) Magnified view of the superimposed acetylated lysine-binding pocket of MOZ (blue) and DPF3b (salmon), with key interacting residues labeled. (G) In vitro peptide pull-down assays like that in Figure 2H. (G) Sequence alignment of tandem PHD fingers of MOZ and DPF3b. Sequence numbers are shown at the right of the corresponding protein sequences. Highly conserved residues in the PHD fingers are highlighted with a purple background. Zinc-coordinated cysteine and histidine residues are grouped by a gray background. Residues responsible for interacting with K14ac, K4me3, and unmodified R2 are highlighted in yellow, red, and orange, respectively. Stars mark residues subject to the mutational analysis in this study.
acetyl amide of H3K14ac forming a pair of hydrogen bonds to the carboxyl oxygen of Asp263 and the sulfur of Cys313, whereas the backbone酰amide oxygen of Lys14 is involved in another hydrogen bond with the Arg289 side chain. Although the Asp263 and Arg289 at the hole of the binding pocket in DPFB3 change to Ser210 and an Asn235 in MOZ [Fig. 3G], water-mediated hydrogen bonds between the carboxyl oxygen of the acetate and Ser210 hydroxyl oxygen as well as Asn235 amino nitrogen were observed in the complex structure [Fig. 3E]. These interactions stabilize the acetate group and indicate the possibility of the acetyl amide of H3K14ac forming hydrogen bonds with Ser210 and Asn235 in a similar manner. Supporting this, alanine substitution of Ser210 or Asn235 causes a reduction in the binding affinity of PHD12 to the H3K14ac peptide [Fig. 2G], and the double mutation destroys the K14ac binding almost entirely [Fig. 3F]. Conversely, we also designed a double point mutation of S210D and N235R on MOZ, interested in the binding properties when these two key residues are identical to the corresponding residues of DPFB3. A further ITC-binding assay showed an almost unaffected K14ac binding of this mutant [Figs. 2G, 3F], indicating that S210 and N235 in MOZ are very likely to suffice for K14ac binding compared with their aspartate and arginine counterparts in DPFB3. In summary, these results indicate that MOZ recognizes the acetylated Lys14 by a hydrophobic pocket and polar residues in PHD1, which is similar to that in DPFB3.

Besides the K14ac-binding site, there are two major differences between the tandem PHD fingers of MOZ and DPFB3 from a global aspect. The first is that the C-terminal α helix in DPFB3 PHD2 changes to a short loop owing to a proline residue. The second is that the loop ahead of the first β strand of MOZ PHD2 is longer than that of DPFB3 PHD2 [Fig. 3D], making it possible to accommodate the trimethylation of Lys4 on histone H3 [Fig. 1C; Supplemental Fig. S3].

Aiming to determine the contribution of MOZ PHD1 to the histone H3 binding directly, we analyzed peptides comprising only the portion of the H3 tail that binds to PHD1 with and without acetylated K14, as has been mentioned before. As shown in Figure 1C and Supplemental Figure S6, however, histone H3 peptides do not bind to PHD12 without the N-terminal eight residues; even Lys14 was acetylated, indicating the indispensable role of the N-terminal residues of H3 peptides in the binding process. In fact, as methylation and alanine mutation on R2 abolish binding completely [Fig. 1E], it is very unlikely for PHD12 to show any affinity to an H3 peptide construct like unH3 (9–18) without unmodified R2.

Considering the weak binding of acetylated Lys14, we were curious which residues on H3 around K14 were also involved to guarantee the Lys14 specificity. The structural comparison with the DPFB3–H3K14ac complex indicated T11 on histone H3 as the only residue with the potential to also be involved in binding [Fig. 3D]. Therefore, we designed the T11A mutation on histone H3 peptide and then measured its binding affinity to PHD12 by ITC and NMR. The ITC data showed a reduction in overall binding, with the $K_d$ value rising from 23.3 ± 2.5 μM to 55.9 ± 9.5 μM [Fig. 1C]. In addition, K243 demonstrated an apparent discrepancy in chemical shift changes upon H3 and H3T11A binding, indicating its role in tethering T11 to histone H3 and therefore optimizing the binding environment of acetylated K14 [Fig. 3D, right]. By combining the NMR titration results with the ITC data, we could conclude that the residues surrounding K14 may contribute to the H3 peptide binding to PHD12, which points to a scenario where instead of recognizing acetylated lysine per se, PHD12 is capable of selecting K14ac in a sequence-specific manner.

**PHD12 binding to histone does not affect the HAT activity of MOZ in vitro**

MOZ [as a HAT], its HAT activity, and its substrate specificity are under strict regulation by other components of the MOZ/MORF complex. Both BRPF1 and ING5 have been shown to up-regulate its acetyltransferase activity [Ullah et al. 2008], and the presence of full-length ING5 was further reported to result in preferential acetylation of methylated H3K4 histone peptides [Champagne et al. 2008]. In another report, the PHD finger ahead of a demethylated Jumonji domain in PHF8 not only directs the substrate specificity of PHF8, but also increases its enzymatic activity by 12-fold [Horton et al. 2010]. Since PHD12 is the only histone recognition module in MOZ, whether its interaction with histone H3 also affects the HAT activity is an interesting question. To address this issue, a truncated MOZ fragment comprising the MYST domain, which is responsible for HAT activity, PHD12-linker-MYST [Fig. 1A]; mutant PHD12 (D282A)-linker-MYST; and a PHD12 control were constructed with a C-terminal His tag and expressed in *Escherichia coli*, respectively. After affinity purification, their enzymatic activities were monitored based on an 3H-based radioactive HAT assay using excessive recombinant histone octamers as substrate. However, although the instability of the PHD12-linker-MYST protein led to long error bars, MYST, wild-type PHD12-linker-MYST, and the mutant all exhibited similar HAT activities against all of the histones compared with the PHD12 control [Supplemental Fig. S7A,B]. By the same token, with the D282A mutant, a HAT assay using recombinant histones containing a H3R2A mutation as substrate was performed. Compared with the wild-type histone octamers, the mutant substrate hardly showed any difference in HAT activity [Supplemental Fig. S7C,D]. All of the above experiments combined indicated that the HAT activity is independent of PHD2 binding to histone H3 at the histone octamer level. The same conclusion could also be made from the competition experiment, in which the excessive H3 [1–18K14ac peptide was added to the recombinant histone octamers while its HAT activity was monitored [Supplemental Fig. S7E]. As the ITC data revealed no direct interdomain physical contact between PHD12 and the MYST domain [Supplemental Fig. S8], it is likely that PHD12 would never directly affect the enzymatic activity of the MYST domain.

Histone octamers might represent a poor substrate to study whether the potential association of PHD fingers...
with acetylated histones could influence HAT activity because we cannot rule out the possibility that these two adjacent domains work on different nucleosomes. In order to better represent its native substrate, we performed the HAT assay with oligonucleosomes extracted from HeLa cells, which are modified and thus contain the ideal binding motif for the PHD fingers. However, we failed to detect any HAT activities of either PHD12-linker-MYST or the MYST domain only [Supplemental Fig. S7F]. A probable explanation rests on the requirement of whole-complex integrity, which is consistent with the previous report that MOZ/MORF HAT activity against oligonucleosomes needs BRPF1 and ING5 [Ullah et al. 2008].

**PHD12 associates with histone H3K14ac in vivo**

To assess the interaction of MOZ PHD12 with histone H3K14ac in vivo, a coimmunoprecipitation assay was performed. Lysates from 293T cells expressing Flag-PHD12 were immunoprecipitated with anti-Flag M2 affinity gel followed by Western blot analysis with anti-H3 and anti-H3K14ac antibodies, respectively. Compared with the plasmid control, Flag-PHD12 markedly bound to nucleosomes with H3K14ac modifications [Fig. 4A], indicating that MOZ PHD12 physically associates with the H3K14ac mark in vivo. Notably, the enrichment of acetylated H3K14 relative to the input is obviously less than that of the total H3, indicating that not all histone H3 in PHD12-bound nucleosomes is acetylated at Lys14. As nucleosomes, which contain two copies of histone H3, most probably remained intact during our experiment procedures, these K14ac-free histone H3 proteins might interact with PHD12 through unmodified R2, although this speculation needs to be further confirmed. Furthermore, confocal fluorescence imaging demonstrated that the ectopically expressed Flag-MOZ fusion protein is colocalized with H3K14ac sites in the H1299 cell nucleus, which became much less apparent in the case of the acetyl-lysine-binding-deficient mutant S210A [Fig. 4B]. This could be additional evidence of the physiological association of MOZ PHD12 with H3K14ac in vivo.

**PHD12 contributes to the localization of MOZ on the HOXA9 promoter**

Previous studies in mice and zebrafish indicated the key role of MOZ in regulating HOX gene expression [Crump et al. 2006; Voss et al. 2009]. Moreover, one recent study in HEK293T and CD44+ cells has demonstrated the binding of MOZ to the HOXA9 gene promoter [Paggetti et al. 2010]. In light of our findings that the tandem PHD finger of MOZ could interact with the histone H3 tail, an interesting question emerges as to whether this binding contributes to a gene-specific targeting. Therefore, we carried out ChIP assays followed by quantitative PCR (ChIP-qPCR) using 293T cells with transient expression of MOZ in full length. As shown in Figure 5, A and B, exogenous MOZ was efficiently recruited to the promoter of HOXA9, but not to the promoter of HOXB1, HOXC5, and HOXD12.

Based on our structural information, we further tested the contributions of each domain of PHD12 in this targeting process by introducing two point mutations, S210A and D282A, as mentioned above, which disrupts its recognition of acetylated K14 and unmodified R2 on histone H3, respectively. The results showed that without affecting the expression level [Fig. 5C] and the nuclear localization of MOZ [Supplemental Fig. S9], the mutant S210A or D282A has a destructive effect on the localization of MOZ on the HOXA9 promoter by a reduction of >50% [Fig. 5B,D]. In these assays, the constant low levels of target gene in groups employing normal mouse IgG as the input control [Fig. 5C] ensured the specificity of immunoprecipitated MOZ to the promoter of HOXA9.

**MOZ PHD12 binding H3 is important for the HOXA9 promoter acetylation and HOXA9 transcription**

MOZ (as a HAT) is responsible for H3K9 or K14 acetylation [Doyon et al. 2006; Voss et al. 2009]. As PHD12 functions to target MOZ to the HOXA9 promoter, it is conceivable to speculate that this PHD12-dependent enrichment could lead to enhanced promoter acetylation. To address this issue, we monitored the acetylation level of the HOXA9 promoter region by ChIP using antibodies against acetylated H3K9 and acetylated H3K14 followed by qPCR. Considering our previous failure in detecting any HAT activity against HeLa oligonucleosomes using a MOZ fragment alone in the absence of a complex and the reports that BRPF1 could greatly enhance its HAT activity [Doyon et al. 2006], we cotransfected GFP-BRPF1 with Flag-MOZ [wild type or mutants] into 293T cells to better preserve the enzymatic activity of the ectopically expressed MOZ. As shown in Figure 6A, 293T cells expressing wild-type MOZ, but not cells expressing the S210A or D282A mutant, exhibited a more than twofold increase in the levels of H3K14 acetylation on the HOXA9 promoter compared with control cells. Notably, the levels of H3K9 acetylation

**Figure 4.** MOZ associates with histone H3K14ac in vivo. [A] Lysates from 293T cells expressing Flag-tagged PHD12 or control vector were immunoprecipitated by anti-Flag M2 affinity gel followed by Western blot analysis with the indicated antibodies. [B] Colocalization of endogenous H3K14ac with ectopically expressed MOZ or its acetyl-lysine-binding-deficient mutant S210A was examined by immunofluorescence assay with the indicated antibodies in H1299 cells.
on the HOXA9 promoter remained similar in all of the cells expressing wild-type or mutant MOZ. These results indicate the H3K14-specific acetylation role of MOZ at the HOXA9 gene promoter. Combined with the previous ChIP data, these results provide a promising picture where the interaction between PHD12 and histone H3 helps MOZ promote the HOXA9 promoter acetylation, mainly at the K14 site. Meanwhile, consistent with previous reports (Voss et al. 2009; Mishima et al. 2011), we found that the global acetylation level remained unperturbed in the absence or presence of cotransfected BRPF1 (Fig. 6B), indicating that the enhancement of the acetylation level might be restricted to specific genes.

Generally, acetylation level is positively correlated with gene transcription. As the interaction between PHD12 and H3 is important for the HOXA9 promoter acetylation, we next examined whether this interaction functions in the HOXA9 gene transcription. As shown in Figure 6C, with the expression of wild-type MOZ, but not the S210A or S282A mutant, HOXA9 mRNA levels were significantly elevated by more than onefold. Based on these results, we could conclude that the interaction between MOZ PHD12 and histone H3 is important for the HOXA9 gene transcription.

On the other hand, under the condition of cotransfection of GFP-BRPF1 and Flag-MOZ (wild type or the S210 or D282 mutant), we analyzed the distribution of BRPF1 and MOZ by fluorescence colocalization assays. Consistent with the previous reports (Laue et al. 2008; Ullah et al. 2008), the overlap of the fluorescence signal from GFP and immune probes against MOZ indicates that the spatial colocalization between MOZ and BRPF1 was unaffected by the mutations (Supplemental Fig. S10). As BRPF1 is shown to be the scaffold of the tetrameric complex and sufficient for the complex formation in vivo (Ullah et al. 2008), it is promising that the biological consequence that we witnessed in this study is largely under its native condition as a complex.

Overall, consistent with our in vitro findings, we identified the tandem PHD finger as a H3 unmodified R2 and acetylated K14 reader with a regulatory function in HOXA9 promoter acetylation and hence the gene transcription. Combined with our previous data, a mechanism could be proposed that through the recognition of the H3R2me0–H3K14ac mark by PHD12, MOZ could be guided onto the HOXA9 locus, where it acetylates adjacent histone and finally switches the HOXA9 gene to its active state.

Discussion

The tandem PHD finger is a functional unit

Extensive studies indicate that histone acetylation is an important epigenetic mark for gene transcription in a chro-
matin context. Knowledge concerning effector modules capable of recognizing acetyl-lysine has long been limited to bromodomains. The tandem PHD finger of DPF3b, as a novel acetyl-lysine-binding module, was identified recently (Lange et al. 2008; Zeng et al. 2010). Due to the current limited information of the tandem PHD finger, it is necessary to further investigate its potential and underlying structural mechanism for specific recognition of acetyl-lysine. Our results show that the tandem PHD finger of MOZ is a histone recognition module that combinatorially reads unmodified histone H3R2 and acetylated H3K14. As described above, the two PHD fingers associate with each other intimately through numerous interactions, which define the two PHD fingers as an integral structural module.

Several nuclear proteins—such as DPF1/2/3b, MLL3, and PHF10—containing the tandem PHD module were identified by the BLAST program using the PHD12 sequence as a query. The sequence alignment indicates that all individual PHD fingers in the modules adopt a canonical Cys4-His-Cys3 motif to coordinate zinc atoms. The two PHD fingers are connected by a short two-residue linker and form an integral module [Supplemental Fig. S11]. Residues that are involved in R2 and K4 recognition by the second PHD finger are conserved, except for the substitution of acid residue by alanine in MOZ, which might lead to the accommodation for methylation of K4. For acetylated K14 binding, a hydrophobic pocket comprised of three residues in the first PHD finger was conserved among all of these proteins; however, the residues responsible for hydrogen bond formation with the acetyl amide and the backbone acylamide oxygen of acetylated H3K14 are variable. In PHF10 and MLL3, the residues corresponding to Ser210 in MOZ change to hydrophobic residues, which might decrease or prevent their binding to acetylated lysine, indicating an unconserved role of the first PHD finger in histone recognition. Notably, both MLL3 and MOZ are histone-modifying enzymes responsible for H3K4me and H4K14ac, respectively. Generally speaking, PHD fingers in this kind of protein regulate their chromatin localization and activity in a highly dynamic manner. In fact, tight binding of PHD fingers to histone might compromise the dynamic regulations by fixing the enzymes to chromatin, providing a possible interpretation of the weak binding of the tandem PHD fingers of MOZ and MLL3 to histone. Therefore, as a partially conserved functional unit, the tandem PHD finger deciphers the histone code to modifying enzymes, allowing them to function in a dynamic process.

Figure 6. MOZ PHD12 binding of H3 is important for the HOXA9 promoter acetylation and HOXA9 transcription. (A) Lysates from 293T cells expressing wild-type MOZ or the S210A or D282A mutant were subjected to ChIP assays with anti-K9ac and anti-K14ac antibodies, respectively, followed by qPCR analysis with HOXA9 promoter-specific primers. The data are represented as mean ± SD of three real-time PCR replicates. (B) The global acetylation level of the cells in the absence or presence of cotransfected BRPF1, which was detected by Western blot analysis with the indicated antibodies. (C) The relative mRNA levels in cells cotransfected with MOZ (wild type or mutant) and BRPF1, which were normalized by β-actins. The data are represented as mean ± SD of three real-time PCR replicates.
Unique recognition of unmodified histone H3K4 (H3K4me0) by PHD2 of MOZ

Previous studies have identified a group of PHD fingers characterized by binding to H3K4me0, including those of a BRAF35–HDAC complex protein [BHC80], BRPF2, autoimmune regulator [AIR], DNA (cytosine-5)-methyltransferase 3-like protein [DNMT3L], and DPF3b. The interactions underlying this type of recognition are generally mediated by direct hydrogen bonds between the oxygen atoms on a single or couple of conserved residues and the amino hydrogen on Lys4 (Supplemental Fig. S12). These conserved residues contain negatively charged atoms in all cases, such as D498 and the backbone oxygen of E497 in BHC80, D212 and D214 in BRPF2, N295 and D297 in AIR, D88 and D90 in DNMT3L, and E315 and D328 in DPF3b. In our case, however, this recognition is indirect, resorting to a water molecule for a tetrahedral hydrogen bond network involving two oxygen atoms in E261, one oxygen atom in N274, and an amino hydrogen in K4 on H3 (Fig. 2E; Supplemental Fig. S12). The inserted water molecule sheds light on a novel H3K4me0 recognition pattern, the introduction of which disrupts the direct polar contacts between PHD finger and H3K4me0 therefore allows relatively weak interactions. As a result, unlike previously identified H3K4me0-binding PHD fingers whose recognition would be abrogated totally by the methylation of H3K4, the tandem PHD finger in MOZ tolerates even the trimethylation on H3K4me3 (Champagne et al. 2008). Based on recent studies and our current work, we proposed a hypothesis model about combinatorial actions of epigenetic readers in the MOZ complex. Like that in the HBO1 [Saksouk et al. 2009] and NuA3 [Taverna et al. 2006] complex, the PHD finger of ING5 binds histone H3K4me3 to recruit the MOZ complex to the target chromatin sites (Fig. 7A; Champagne et al. 2008). After that, the first PHD finger of the BRPF protein binds H3K4me0 [Qin et al. 2011], and the tandem PHD finger of MOZ binds to unmodified H3R2, stabilizing the complex to perform acetyl transfer reactions against histone H3 (Fig. 7B). Accompanied by chromatin relaxation, the positive feedback mechanism begins to function, spreads the H3K14ac mark around, and finally switches the target gene to its active state (Fig. 7C). Supportive of these findings, another study showed that MOZ and MLL, an H3K4me2/3 HMT complex, interact, colocalize, and functionally cooperate on multiple HOX promoters [Paggetti et al. 2010], highlighting the relationships between MOZ and the modifications of H3K4me3 and H3K14ac. Notably, methylation of H3R2, like a molecular switch, abrogates the trimethylation of H3K4 by the Set1 methyltransferase [Guccione et al. 2007; Kirmizis et al. 2007] and functions as a negative regulator of N-terminal H3 tail binding [Iberg et al. 2008]. In our case, dimethylation of H3R2 greatly impedes the interactions between the MOZ complex and histone H3 by abolishing the bindings of PHD fingers of MOZ and ING5 to N-terminal H3 tails. However, more details regarding the H3R2me2-mediated regulation of MOZ and histone H3 acetylation depend on future investigations.
on a single histone tail and beyond may have a significant, if not dominant, role in chromatin transactions.

Materials and methods

Expression, purification, and isotope labeling of MOZ PHD12

The DNA fragments of human MOZ tandem PHD finger residues 204–313 were amplified from the human brain cDNA library (Clontech) by PCR reactions. Then, the DNA fragments were ligated into the NdeI/XhoI-cleaved plasmid pET-28a(+) (Novagen), which was modified with thrombin cleavage site deletion or substitution of a TEV cleavage site, generating fusion proteins with a 6xHis tag at the N terminus with or without a TEV cleavage site. MOZ PHD12 mutants were generated by the conventional PCR method using a pET-28a(+)–PHD12 plasmid as template. All of the recombinant MOZ PHD12 proteins were produced in *E. coli* BL21 (DE3). Cells were grown in either LB medium for unlabeled samples or minimal medium supplemented as appropriate with glucose or [13C6] glucose and [15NH4]Cl for 15N and 13C samples, and 0.1 mM ZnSO4 was added. Generally, the protein expression was induced at A600 = 0.8–1.0 with 0.5 mM IPTG for 24 h at 16°C. The expressed recombinant protein was purified using a Ni-chelating column (Qiagen) followed by TEV cleavage (for crystallography only) and Superdex 75 column gel filtration. The purified protein was exchanged to buffer A (20 mM HEPES, 100 mM NaCl at pH 8.0) as an NMR sample.

Peptide synthesis

H3 [1–18]K14ac, and H4 [11–21]K16ac were used in this study, their sequences are listed in Supplemental Table S3, and they were synthesized by GL Biochem [Shanghai]. Biotinylated peptides H3 [1–18], H3 [1–18]R2me2a, H3 [1–18]R2me2s, and H3 [1–18]K14ac were synthesized by Scilight Biotechnology [Beijing]. They were supplied with stringent analytical specifications [purity level >98%], which included HPLC and MS analysis. Before use, peptides were weighed and dissolved in buffer A, and the pH of each sample was systematically controlled and, if necessary, adjusted to a value of 8.0 by addition of diluted NaOH or a HCl solution.

**NMR spectroscopy**

All spectra were recorded at 25°C on a Bruker DMX500 or DMX600 spectrometer. Both the purified 15N-labeled and 13C, 13N-labeled proteins were dissolved to a final concentration of 0.8 mM in 500 μL of buffer A with 10% D2O. All NMR experiments were performed at 293 K on a Bruker DMX500 or DMX600 spectrometer.

The following spectra were recorded to obtain backbone and side chain resonance assignments [Clare and Gronenborn 1994]: two-dimensional (2D) 1H-15N HSQC; three-dimensional (3D) triple-resonance spectra HNCO, HNCACB, CBCA(CO)NH, CBCANH, HBBCHA(CO)NH, C(CO)NH-TOCSY, and H(CCO)NH-TOCSY; 2D 1H–13C HSQC; and 3D HCCH-TOCSY and HCCH-CBCANH, HBHA(CO)NH, C(CO)NH-TOCSY, and H(CCO)NH-TOCSY. NOE (nuclear Overhauser effect) distance restraints were generated using 15N-edited NOEASY and 13C-edited NOEASY spectra using 110-msec mixing times. NMR data were processed by NMRPipe and NMRDraw software [Delaglio et al. 1995] and were assigned with Sparky (T.D. Goddard and D.G. Kneller, University of California at San Francisco). The Talos program [Shen et al. 2009] was used to obtain the backbone dihedral angles (Φ and ψ) in secondary structures on the basis of chemical shift information. Hydrogen bond restraints were defined from slow-exchanging amide protons identified after exchange of the H2O buffer to D2O.

**Structure calculations**

Structure calculation for PHD12 was performed on the basis of proton–proton NOE restraints and dihedral angle (Φ and ψ) restraints with a simulated annealing protocol using the CNS version 1.2 program [Brunger et al. 1998]. High-temperature torsion angle dynamics were performed for 15 psec at 50,000 K [1000 steps], followed by a 15-psec cooling phase. Initial structure calculations included only hydrogen bonds in defined secondary structures from Talos. In the following refinement calculation, only hydrogen bonds whose donors could be identified unambiguously were added. In the final calculations, an ensemble of 200 structures [with no distance violations >0.3 Å and no dihedral angle violations >5°] was generated from unambiguous NOEs previously determined. Twenty models were selected on the basis of energetic criteria [low total energy, using the accept.inp routine] to form a representative ensemble of the calculated structures.

**Chemical shift perturbation**

For detecting the histone peptides with several modification sites on MOZ PHD12, 0.3 mM 15N-labeled PHD12 was used. After the 1H-15N HSQC spectrum of free MOZ PHD12 was recorded, the sample was titrated with histone peptides by the method described by Weber et al. [1991]. All HSQC spectra for mapping histone peptide-binding interfaces on MOZ PHD12 were performed on a 600-MHz Bruker at 298 K.

**FPAs**

FPAs were performed in buffer B [20 mM HEPES, 150 mM NaCl at pH 7.0] at 293 K using a SpectraMax M5 microplate reader system [Molecular Devices]. The wavelengths of fluorescence excitation and emission were 485 nm and 522 nm, respectively. Each well-plate contained 100 nM C-terminal fluorescein isothiocyanate (FITC)-labeled H5K14ac peptide [residues 1–18] probe and different amounts of PHD12 or mutants [concentrations from 0 μM to ~100 μM] with a final volume of 200 μL. For each assay, peptide-free controls (PHD12 or mutants only) were included.

**Peptide pull-down assays**

Biotinylated histone peptides (1 μg each) that were used for the pull-down assays were incubated with 2–5 μg of purified GST-PHD12 for 2 h at room temperature in binding buffer [20 mM Tris-HCl at pH 7.5, 150 mM NaCl]. After 1 h of incubation, Streptavidin beads [ThermoScientific] were washed three times using binding buffer with 0.1% Triton X-100 and were subjected to Coomassie blue staining.

**X-ray crystallography**

The 6xHis tag of the recombinant MOZ PHD12 protein was removed by TEV enzyme digestion. Then, the protein was purified by Superdex 75 column gel filtration and monoQ anion exchange. The MOZ PHD12 complexes with H3K14ac were prepared by mixing 8.5 mg of PHD12 with 3 mg of H3K14ac [1–18] peptide in a 1-mL volume and incubated for a further 15 min on ice. Crystals were grown at 20°C by the sessile droplet vapor diffusion method by mixing 1 μL of complex with 1 μL of reservoir solution. Crystals for the complex were produced from a reservoir solution containing 0.2 M ammonium acetate, 0.1 M tri-sodium citrate, and 30% PEG4000 [pH 5.6].

A SAD data set was collected on beam line 17U at Shanghai Synchrotron Radiation Facilities (SSRF) at a wavelength of 0.97 Å, and the data were processed using programs in CCP4 package [Collaborative Computational Project, Number 4 1994]. All of the four expected endogenous zinc atom positions were determined with SHELX C/D [Sheldrick 2008], and the initial phases of calculations and density modification were produced with SHELX E [Sheldrick 2008]. An initial model was automatically built by the program ARP/wARP [Langer et al. 2008]. The model was further built and refined using COOT [Emsley et al. 2010] and Refmac5 [Murshudov et al. 1997], respectively.

**HAT activity assay**

The HAT activity of the recombinant PHD-MYST domain and MYST domain were determined using a 3H-based radioactive HAT assay as described [Holbert et al. 2007]. The substrate used in the HAT assay was histone octamer reconstituted as described by Luger et al. [1999]. The following reaction protocol was applied for 30-μL reaction volumes: Two concentrations of MOZ PHD-MYST and its mutants [30 or 90 nM] were incubated with 1 μg of histone octamer and 0.25 μCi of Ac-CoA in HAT buffer [20 mM Tris-HCl at pH 7.6, 30 mM sodium chloride, 5% [w/v] glycerol, 10 mM sodium butyrate, 1 mM phenylmethylsulfonyl fluoride] for 30 min at 30°C. Buffer control reactions not including enzyme were also performed. After incubation, the reactions were stopped by adding 5 × SDS sample buffer, separated using SDS-PAGE, and then transferred onto a PVDF membrane. The transferred protein samples were stained by Coomassie brilliant blue R-250, processed by EN3HHANCE spray, and subsequently exposed to
Amersham HyperfilmTM MP for 24 h at –80°C. The bands on the PVDF membrane corresponding to histone octamer could be subject to scintillation counting after the exposure. For HAT assays of different fragments of MOZ, the concentrations of MOZ fragments and histone octamer were switched to 200 nM and 2 μg per reaction, respectively.

Coimmunoprecipitation and Western blot

Two 6-cm dishes of 293T cells (transiently transfected with control Flag plasmid or Flag-PHD12 plasmid) were harvested, washed with PBS, and ultrasonicated in a cold PBS buffer containing protease inhibitor cocktail. After centrifugation, 1 mg of total proteins was then incubated with anti-Flag M2 affinity gel (Sigma) for immunoprecipitating proteins with gentle shaking overnight at 4°C. Immunoprecipitation complexes were washed twice in ice-cold PBS buffer. Proteins were eluted by boiling in Laemmli buffer. Eluted proteins were then separated by SDS-PAGE and electroblotted to nitrocellulose membranes. Membranes were blocked in 1× PBS-T (0.1%) and fat-free dry milk or bovine serum albumin (BSA; 5%) [blocking buffer] for 1 h at room temperature. Membranes were incubated with the primary antibodies diluted in the blocking buffer overnight at 4°C. Membranes were washed three times in 1× PBS-T (0.1%) during 10 min each. Secondary antibodies conjugated with horseradish peroxidase were added, and the membranes were incubated for 1 h at room temperature. Membranes were washed three times in 1× PBS-T (0.1%) for 10 min each. ECL Western blotting reagent kit (ThermoScientific) was used for protein detection.

Immunostaining and fluorescence colocalization

Cells were cultured in 24-well chamber slides. After washing once with PBS, cells were fixed in 4% paraformaldehyde and permeabilized in PBS containing 0.5% Triton X-100 for 5 min at room temperature. After blocking with 5% BSA in PBS for 1 h, cells were incubated with primary antibodies and donkey anti-mouse immunoglobulin G conjugated with rhodamine antibody (Jackson ImmunoResearch) or goat anti-rabbit immunoglobulin G conjugated with FITC antibody (Jackson ImmunoResearch), respectively. Hoechst 33,342 (Sigma) was used for nuclear staining. After extensive washing, cells were analyzed by an Olympus IX71 inverted microscope coupled to an Olympus DP70 high-resolution color camera or by a Zeiss LSM710 laser confocal system.

ChIP

293T cells were cultured in DMEM medium supplemented with 10% fetal bovine serum (Gibco). Transient transfection was performed using EZ ChIP kit (Millipore). Briefly, 293T cells (1 × 107 cells) were fixed with 1% formaldehyde and then neutralized by adding 0.125 M glycine. Cells were collected and lysed in cell lysis buffer containing SDS and cocktail protease inhibitor. The lysate was sonicated to obtain soluble chromatin with an average length of 1000 bp. After 1:10 dilution in dilution buffer, chromatin solutions were precleared and then incubated with IgG, anti-Flag antibody (Sigma), anti-H3 antibody (Abcam), anti-H3K9ac antibody (Millipore), or anti-H3K14ac antibody (Millipore), respectively. Next, the mixture was incubated overnight at 4°C on a rotating platform. The immunocomplexes were captured by protein G-Sepharose beads. After extensive washing, the bound DNA fragments were eluted, and the resulting DNA was subjected to real-time qPCR analysis using the primer pairs shown in Supplemental Table S3.

MOZ PHD12 is a H3R2- and H3K14ac-binding module

Accession numbers

The atomic coordinates and structure factors have been deposited in the PDB with the following accession codes: PHD12 in the free state (2LN0) and in complex with H3 [1–18]K14ac peptide (3V43). The Biological Magnetic Resonance Data Bank (BMRB) accession code for the NMR structure of PHD12 is 18142.

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MOZ PHD12 is a H3R2- and H3K14ac-binding module


