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Substrate- and Cofactor-independent Inhibition of Histone Demethylase KDM4C

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Supporting Information

ABSTRACT: Inhibition of histone demethylases has within recent years advanced into a new strategy for treating cancer and other diseases. Targeting specific histone demethylases can be challenging, as the active sites of KDM1A-B and KDM4A-D histone demethylases are highly conserved. Most inhibitors developed up-to-date target either the cofactor- or substrate-binding sites of these enzymes, resulting in a lack of selectivity and off-target effects. This study describes the discovery of the first peptide-based inhibitors of KDM4 histone demethylases that do not share the histone peptide sequence or inhibit through substrate competition. Through screening of DNA-encoded peptide libraries against KDM1 and -4 histone demethylases by phage display, two cyclic peptides targeting the histone demethylase KDM4C were identified and developed as inhibitors by amino acid replacement, truncation, and chemical modifications. Hydrogen/deuterium exchange mass spectrometry revealed that the peptide-based inhibitors target KDM4C through substrate-independent interactions located on the surface remote from the active site within less conserved regions of KDM4C. The sites discovered in this study provide a new approach of targeting KDM4C through substrate- and cofactor-independent interactions and may be further explored to develop potent selective inhibitors and biological probes for the KDM4 family.

The dynamic regulation of gene expression is controlled by a range of mechanisms, among which reversible posttranslational modifications (PTM) of the N-terminal tails of histone proteins play an important role through affecting chromatin structure.1,2 Deregulation of histone-modifying enzymes has been shown in a number of diseases, including cancer;3 thus, inhibitors of histone-modifying enzymes are interesting probes for investigating the biological role of these enzymes and their potential as therapeutic targets.

Until the discovery of the histone demethylase KDM1A in 2004, histone methylation was thought to be an irreversible epigenetic mark.4 KDM1A and its paralog KDM1B are FAD-dependent amino oxides demethylating mono- and dimethylated lysine 4 on histone H3 (H3K4me2/me1). The KDM4 family of Jumonji-domain containing demethylases was identified in 20065 and consists of the six members KDM4A, -B, -C, -D, -E, and -F. Among those, KDM4E and -F are considered pseudo-genes.6,7 while KDM4A-D produce enzymatically active gene products. KDM4A-D are known to demethylate H3K9me2/3, H3K36me3/2, and H1K26me3/2 through a hydroxylation reaction requiring the cofactors Fe(II) and 2-oxoglutarate (2-OG).5,9 Due to their elevated activity and expression in several forms of cancer, KDM1 and -4 proteins are recognized as oncogenes.10,11 KDM1 is, among others, associated with prostate, bladder, and estrogen-receptor-negative breast cancer.12 The KDM4 family of histone demethylases has repeatedly been shown to be involved in progression of hormone dependent cancers, such as breast and prostate cancer through coregulating hormone receptors.13–15 The development of selective KDM1 or -4 inhibitors is impeded by the high structure and sequence conservation of these enzymes; most known inhibitors mimic either the FAD cofactor (KDM1) or 2-OG through Fe(II)-binding (KDM4) and hence interact with a plethora of other targets, such as 2-OG-dependent oxygenases through iron chelation.16–19 The conjugation of iron chelating compounds to the truncated histone peptide substrate has been investigated as well,20,21 and it resulted in the discovery of the first KDM4 selective inhibitors. However, there is a need for additional strategies to target histone demethylases through noncofactor and nonsubstrate interactions. Novel inhibitory
scaffolds targeting alternative sites on histone demethylases are warranted, as they might hold the key to subfamily and isoform selectivity.

Herein, we present the discovery of several peptide binders of the histone demethylases KDM1A, -4A and -4C using phage display, which are not related to the sequence of their natural histone peptide substrates. Two of these peptides were developed into inhibitors of KDM4C by amino acid replacement, truncation, and chemical modifications. The inhibitors were found to target KDM4C via substrate-independent interactions on the surface of the enzyme located in neighboring regions of the highly conserved active site and within less conserved regions.

**RESULTS AND DISCUSSION**

**Phage Display Screening.** Phage display screening is a versatile tool for the discovery of peptides binding to biological targets such as proteins. A phage library displaying random peptide sequences fused to the N-terminus of the phage protein pIII was screened against the catalytic domains of histone demethylases KDM1A, -4A, -4B, -4C, -4D, and -4E. The library consisted of linear 7- and 12-mer peptide sequences (X7-pIII was screened against the surface-immobilized target proteins, phages binding to KDM1 and -4 proteins were amplified, and the KDM1 and -4 binding phages did not share any sequence motifs. The apparent affinity of the phages was increased by only ∼10-fold. Truncation of 2 from the N- or both termini, leading to peptides 15 and 17, resulted in loss of inhibitory activity, while truncation from the C-terminus (16) reduced inhibitory potency by ∼3-fold. Interestingly, also N-terminal acetylation of 2 (24) led to a complete loss of inhibitory activity on KDM4C, indicating that the N-terminal alanine residue serves an important function in inhibiting KDM4C.

**Peptide Development.** In order to increase the inhibitory activity of 1 and 2 which was dramatically decreased in comparison to the peptide-phages, their structures were optimized. It was hypothesized that weak inhibition of 1 in particular, but also 2, might originate from instability of the peptides owing to the presence of aspartate and asparagine residues that are known to destabilize peptides in solution, leading to deamidation and isomerization. Hence, the aspartate residues in peptide 1 were exchanged with alanine and/or glutamate, and the asparagine residues in 2 were exchanged with alanine. Single and double exchange of D6 and D7 to alanine in peptide 1 gave analogues 3–5, and exchange to glutamate gave 6–8. The exchange of D6A (4) led to a dramatic increase in inhibitory activity, and also the double substitution D6E/D7E resulted in an increase in inhibitory potency. These findings indicate that 1 might indeed be inactive due to stability issues. All other analogues of 1 were inactive. For 2, exchanges N6A and N8A gave inactive peptides 9 and 10, whereas the double exchange N6A/N8A yielded the 2-fold less active peptide 11.

The initial synthesis of peptides 1 and 2 included the phage-derived linker sequences, an N-terminal alanine residue, and three C-terminal glycine residues. To reduce the size of the peptides, these nonrandomized sequences beyond the cysteines were removed sequentially. First, peptide 4 was truncated from either the N- or C-terminus, leading to inactive peptides 12 and 13. Truncation from both ends (14) reduced inhibitory activity by only ∼10-fold. Truncation of 2 from the N- or both termini, leading to peptides 15 and 17, resulted in loss of inhibitory activity, while truncation from the C-terminus (16) reduced inhibitory potency by ∼3-fold. Interestingly, also N-terminal acetylation of 2 (24) led to a complete loss of inhibitory activity on KDM4C, indicating that the N-terminal alanine residue serves an important function in inhibiting KDM4C.

**Peptide Sequences Targeting KDM1A, -B, and KDM4A–C Identified from Phage Display, As Well As The Apparent EC50 Values of the Peptide-Phages**

| protein | phage-peptide | sequence | EC50 [μM] |
|---------|---------------|----------|-----------|
| KDM1A   | PP1           | SHFEDWDWGPQGGG | 1.6       |
| KDM1B   | PP2           | AWDVIVDQILQH | 10        |
| KDM4A   | PP3           | GRMDWLGWRELGGG | 11        |
|         | PP4           | SHSMNRAPSLVRGGG | 3.4      |
| KDM4C   | PP5*         | ACKWMDGYYCQGGG | 56       |
|         | PP6*         | ACYTRNNMQCQGGG | 1.3     |

*Peptides cyclized via disulfide bridge on the phage surface.*

identified, and the KDM1 and -4 binding phages did not share any sequence motifs. The apparent affinities of the phages were established to be in the low μM range from competition experiments performed between free and surface-immobilized KDM protein for phage binding (Table 1 and Supporting Information (SI) Figure S1). Cross affinity testing of peptides-phages PP1–6 against all KDM1 and -4 proteins revealed that most phages were specific binders (SI Table S2); however, PP5 displayed some affinity toward KDM1A, and PP3 showed weak affinity for several proteins.

After their identification from phage display, the inhibitory activity of the soluble peptides was assessed on their respective epigenetic target proteins. The peptides were initially synthesized including the phage-derived N- and C-terminal linker sequences, and C-terminally amidated to resemble the fusion to the phage protein pIII. The peptides were tested for demethylase inhibitory activity against their targets KDM1A, KDM1B, KDM4A, and KDM4C (SI Figures S2–6). The two cyclic peptides 1 (derived from PP5) and 2 (derived from PP6) inhibited KDM4C-mediated H3K9me3 demethylation with IC50 values of 1000 μM and 52 μM, respectively (Table 2).
improved inhibition (IC_{50} = 2.6 \mu M), indicating that the more bulky threonine residue could hinder interaction with KDM4C.

Table 2. Amino Acid Sequences of Peptides 1 and 2 and Their Analogues, IC_{50} Values, and Hill Slopes on KDM4C

| peptide sequence | IC_{50} [\mu M]^a | Hill slope |
|------------------|-------------------|-----------|
| 1 ACKWMADGYCGGGb | 1000              |           |
| 2 ACYTRMNQCGGGb  | 52 ± 1.5          | −0.42     |
| 3 ACKWMAMAGCGGGb | >1000             |           |
| 4 ACKWMADGYCGGGb | 8.5 ± 1.5         | −0.56     |
| 5 ACKWMADGYCGGGb | >1000             |           |
| 6 ACKWMADGYCGGGb | ∼520              |           |
| 7 ACKWMADGYCGGGb | >1000             |           |
| 8 ACKWMADGYCGGGb | >1000             |           |
| 9 ACYTRNMQCGGGb  | >1000             |           |
| 10 ACYTRNMQCGGGb | >1000             |           |
| 11 ACYTRMNQCGGGb | 97 ± 1.5          | −0.90     |
| 12 CKWMADGYCGGGb | >1000             |           |
| 13 ACKWMADGYCb   | >1000             |           |
| 14 CKWMADGYCb   | 76 ± 1.3          | −0.93     |
| 15 CYTRNMQCGGGb  | >1000             |           |
| 16 ACYTRNMQCb   | 406 ± 1.3         | −0.55     |
| 17 CYTRNMQCb     | >1000             |           |
| 21 Meta-Xyl-ACYTRNMQCa | 16 ± 1.6 | −0.70     |
| 22 Meta-Xyl-CKWMADGYCb | >1000 |           |
| 23 ACYTRNMQCGGGYGRKKRRQRRb | 0.6 ± 0.02 | −0.46     |
| 24 Ac-ACYTRNMQCGGGb | >1000             |           |

a Determined by LANCEUltra KDM4C Histone H3K9 demethylase assay. \( K_m \) (ccKDM4C) = 220 nM, \( c (H3K9me3) \) = 300 nM. Full dose–response curves are shown in SI Figure S6. b Peptides cyclized by disulfide bridge. c Peptides cyclized by \( m-\alpha-\alpha' \)-dibromoxylene.

Figure 1. Compounds 18–22 cyclized by dibromoxlenes.

Figure 2. Average change in inhibitory activity of alanine analogues of peptide 14 (A) and 21 (B) on KDM4C normalized to the activity of the original peptides. Error bars are standard deviation of the mean (\( n = 3 \)).

Cellular Activity. The effects of compounds 2, 4, and 21 on cellular histone demethylation were investigated through cellular
immunoﬂuorescence assays using osteosarcoma U2OS cells. In addition, the effect of attaching a cell penetrating TAT-peptide to the C-terminus of peptide 2 was investigated (peptide 23). Though an increase in inhibitory activity was observed for 23 (Table 2), no cellular activity on histone demethylase activity could be detected for this or any of the other tested compounds (data not shown). Most likely, the lack of activity was due to low cell-permeability, cellular stability issues, and/or the insufficient inhibitory activity of the compounds.

Interaction with KDM4C. In order to investigate the mechanism by which the peptides inhibit KDM4C in more detail, the two best analogues from each series, 4 and 23, were tested in substrate competition experiments. The Michaelis–Menten kinetics of KDM4C were determined at various inhibitor concentrations, as shown in Figure 3A, B. For both compounds, the $K_m$ value initially decreased by ∼30% upon inhibitor addition but remained constant at higher inhibitor concentrations. In contrast to this behavior, the $V_{max}$ values continuously decrease with increasing inhibitor concentration. These results imply that 4 and 23 do not simply inhibit KDM4C in a substrate-competitive manner, but through a type of noncompetitive inhibition. Also, the shift of the intersection in the Lineweaver–Burk plots to the right of the y-axis (Figure 3C, D) indicates an atypical mechanism of inhibition, potentially through cooperative binding with more than one peptide.

To dissect the mode of interaction between the inhibitors and KDM4C, hydrogen–deuterium exchange (HDX) mass spectrometry (MS) was employed to explore the sites affected by peptide-binding on KDM4C in solution. HDX-MS monitors the HDX of backbone amide hydrogens in a protein by MS; HDX can be localized to different regions of the protein by pepsin proteolysis and mass analysis of the deuterium content of the resulting peptic peptides. A ligand binding event can cause a reduction in the deuterium uptake in regions of the protein involved in binding. Prior to HDX, the peptides were incubated with KDM4C for 30 min at room temperature (RT) to allow

Figure 3. Substrate-dependent inhibition of KDM4C by 4 (A, C) and 23 (B, D). (A, B) The velocity of KDM4C at different inhibitor concentrations is plotted against the substrate concentration, the tables show the respective $K_m$ and $V_{max}$ values. (C, D) Corresponding Lineweaver–Burk plots.

Figure 4. Differential HDX-MS data for KDM4C ± peptide inhibitors. (A) Histograms showing the percent reduction in HDX for KDM4C residues 113–120, 166–174, 205–215, 248–275, 331–344, and 347–356 by peptides 2, 21, 4, and 14. Values are calculated relative to the measured %D value for free KDM4C (n = 3, error bars are standard deviation of the mean). B) HDX data for the four regions of interest are plotted over the structure of KDM4C (pdb 2XML) with 113–120 highlighted in light blue, 166–174 in blue, 248–275 in orange, and 331–344 in light orange. The histone peptide substrate is shown in red.
binding of the peptide to the protein. Upon peptide addition, reductions in deuterium uptake were observed on KDM4C fragments 113–120 and 166–174 for peptides 2, 4, 14, and 21 (Figure 4A and SI Figure S12). Peptides 4 and 14 showed additional alterations on 248–275, and 2, 4, and 14 on 331–344, suggesting distinct binding modes. Inspection of the interacting regions in the X-ray crystal structure of KDM4C indicates two regions affected by peptide binding: the first one spanning residues 113–120 (light blue) and 166–174 (blue) where all inhibitory peptides induce reduced deuterium uptake for KDM4C, and a second one covering 248–275 (orange) and 331–344 (light orange) on KDM4C (Figure 4B). All identified regions are located on the surface of KDM4C and remote from the substrate binding site (red). To verify that no interaction takes place in the active site, regions involved in substrate binding (205–215) and catalytic activity (347–356) were analyzed as well, but no changes in HDX upon peptide addition could be detected (Figure 4A). Thus, inhibition seems to occur through interactions on the surface of KDM4C. The identified sites are located within less conserved regions remote from the active site of KDM4C, suggesting a potential selectivity toward inhibition of distinct isoforms of the KDM4 family. These sites are also remote from the N- and C-terminal tails of the construct and are most likely accessible on the full-length enzyme, since the rest of the protein sequence of the enzyme continues on the back side of the view in Figure 4B. However, it remains to be established that the compounds can inhibit full length enzymes. Considering that the two surfaces (spanning from 113 to 120 and 166–174, and 248–275 and 331–344) are too far apart to be reached by a single peptide molecule, these results support the previously hypothesized cooperative binding of the peptides to KDM4C. Furthermore, the overall negative Hill slopes of the inhibitors (Table 2) also indicate negative cooperative binding of the inhibitors to KDM4C. Interestingly, it has been suggested in the literature that the interactions between phage-display-derived peptides and their target proteins are not only driven by the monomeric sequence of the peptide but by its multimeric presentation on the phage. The results could be interpreted to support this suggestion.

Both the cross affinity screening on phage-level (SI Table S2) and the results from the HDX-MS analysis indicate that the peptides could display a certain level of isoform selectivity. The cross affinity screening showed that PP5 (developed into 4) has the highest affinity on KDM4A, while PP6 (developed into 2 and 21) binds primarily to KDM4C. To test the peptides’ isoform selectivity, 2, 4, and 21 were tested for their inhibitory activity on KDM4A and -B by AlphaScreen, revealing that all three peptides inhibit KDM4A and -B to various degrees, with 21 being the most potent inhibitor, and 2 and 4 showing overall lower activity (SI Figure S9). For all tested compounds, KDM4A shows the highest level of inhibition, followed by KDM4C and then KDM4B. These results are in good agreement with the degree of structural conservation among KDM4A-C, which is highest between KDM4A and -C. The IC50 values of 21 on KDM4A, -B, and -C were determined in separate dose–response experiments from AlphaScreen and ranged from 1.1 μM for KDM4B to 2.1 μM for KDM4A and 3.5 μM for KDM4C (SI Figure S10). These results show that there is no selectivity of the compounds among the KDM4 family. In principle, the compounds could also have activity at other demethylases, but it is the aim of future studies to establish a complete selectivity profile.

Conclusion. In conclusion, phage display screening provided a viable platform to identify inhibitors of histone demethylases. Though the initial hits discovered by phage display showed only moderate inhibitory behavior, it was possible to develop them into more potent compounds through amino acid replacement, truncation, and chemical modifications. The optimized KDM4C inhibitors were analyzed biochemically with respect to their target engagement, suggesting that inhibition takes place through a mechanism that is not competitive and potentially involves cooperative binding. This hypothesis was further supported by the HDX-MS analysis that suggested two distinct binding surfaces on the catalytic core of KDM4C located remote from the active site. Initial optimization of the peptides increased the potency of the inhibitors but did not lead to isoform selectivity. However, since the sites affected by peptide binding were found to be located within a less conserved region of KDM4C, and the phage display also suggested isoform selectivity, the peptide sequences described may be developed into selective inhibitors for individual KDM4 isoforms. It can further be speculated that the peptides target protein–protein interactions, or dimerization sites of KDM4C. Multipeptide complexes containing KDM4C are believed to be important for the recruitment of the demethylase to transcription sites; hence, the peptides could potentially be used as tool compounds to abolish such interactions. The dimerization of KDM4C has been demonstrated in the literature; however, it remains unknown whether the enzyme is functional as a mono- or a dimer. Also, the dimerization interface of KDM4C is unknown, albeit all KDM4 proteins normally crystalize as dimers. Though this dimerization might be artificially induced through the high protein concentrations required for crystal formation, the crystal structure of KDM4C (pdb 2XML) indicates that the dimerization surface is located between the N-terminal JmjN domain and the β-hairpin (residues 112–135). If this site is involved in dimer formation and dimerization is crucial for demethylase activity, the binding of the peptides to this site, as shown by HDX-MS, would disrupt the dimer and inhibit the enzyme. Additional studies are, however, needed to verify this.

Based on our findings, we believe that the herein described compounds are important new lead structures for developing potent, selective tool compounds for both epigenetic therapy and the study of the physiological role of KDM4C. Future studies will aim at increasing potency through developing these interesting scaffolds by further modifications and to further characterize the biological activity of these peptides. Moreover, important goals will be to induce isoform selectivity and cell-activity of the peptides.

### METHODS

**Phage Display.** Two linear 7-mer and 12-mer and a constrained version of the peptide library were used for the selection. Microtiter plates were coated at 4 °C for 16 h with 10 μM purified protein in 1× PBS (20 mM sodium phosphate, 150 mM NaCl, pH 7.4, total volume per well 100 μL). The microtiter plates were subsequently washed with 1× PBS and then blocked with 4% skim milk in 1× PBS for 1 h at RT. Bacteriophages at around 1010–1011 pfu/ml were used for each panning round. The constrained and linear libraries were mixed and panned together in 1× PBS. After incubation for 1 h at RT, the plates were washed 10 times with 1× PBS and bound peptide-phages were eluted with glycine/HCl, pH = 2.2 for 10 min, followed by neutralization with Tris-base, pH = 9.0. Eluted phages were used to infect exponentially growing TG1 cells overnight at 37 °C. The following day, the peptide phages were precipitated from the cell supernatant with phage precipitation buffer (20% (w/v) PEG6000 in 2.5 M NaCl) and redissolved in 300 μL 1× PBS. After 4–5 rounds of panning, single clones were isolated and tested for KDM4A-E and KDM1A and -B.
binding in phage ELISA (procedure described in the Supporting Information). Single stranded DNA was extracted from positive clones, and the DNA was sequenced in the region corresponding to the random peptide region.

More experimental details and the protocols for phage and peptide ELISA are shown in the Supporting Information.

**LANCE Ultra KDM4C Histone H3-Lysine 9 Demethylase Assay.** For inhibition studies on KDM4C, PerkinElmer’s LANCE Ultra JMJ2C Histone H3-Lysine 9 demethylase assay was performed according to the manufacturer’s protocol. The catalytic core domain of KDM4C (ccKDM4C) employed in the LANCE assay was prepared as described previously. Assays were carried out in triplicate measurements at RT; the total reaction volume was 10 μL/well. For inhibition studies, 0.6 fmol ccKDM4C were incubated for 30 min with various concentrations of the peptides prior to addition of the substrate mix, containing 300 nM H3K9me3 peptide substrate. The reaction mixture was allowed to incubate for another 30 min at RT, before the reaction was stopped by addition of the detection mix. The TR-FRET signal was measured under aerobic conditions using a Beckman Instruments DU 2500 microplate reader (Tecan). Reaction mixtures of 200 nM KDM1A, or 480 nM KDM1B) to reaction mixtures were acquired on a Waters SynaptG2 HDMS mass spectrometer. Peptic peptides of KDM4C identified from this analysis is shown in SI Figure S11. The sequence coverage was 91.7% with a redundancy of 1.8%. Mass spectra were processed using the Mass Lynx and DynamX software packages (Waters Corp.). Complete deuteration of control samples was achieved by incubation of 60 pmol KDM4C in the deuterated buffer in the presence of 6 M guanidine for 24 h at 30 °C. Average back exchange (i.e., deuterium loss) was measured as 37% for all samples were compared. The HDX of KDM4C in the presence and absence of peptide into the corresponding deuterated buffer (12.5 mM HEPES pH 7.4, 5 μM FeSO4, 100 μM ascorbate). Nondeuterated controls were prepared by dilution into an identical protiated buffer. All HDX reactions were removed and were immediately frozen and stored at −80 °C. Samples were injected into a cooled Waters nanoAcuity UPLC system for online pepin digestion and rapid desalting of the protein samples. After digestion of the samples, the peptic peptides were trapped on a trap column (Waters VanGuard C18, 1.7 μm, 2.1 × 5 mm) and desalted with 0.68% formic acid in 8% acetonitrile, pH 2.3, 200 μL min−1 for 3 min. Peptides were eluted from the trap to the analytical column (Waters XBridge C18, 1.7 μm, 1.0 × 100 mm) and separated with an 8–40% gradient of 0.68% formic acid in acetonitrile (pH 2.3) over 12 min at a flow rate of 40 μL/min. Positive ion-electrospray ionization mass spectra of eluted peptides were acquired on a Waters SynaptG2 HDMS mass spectrometer. Peptic peptides were identified in separate experiments using collision-induced dissociation tandem mass spectrometry performed with a data-independent (MS3) acquisition scheme. A sequence coverage map of peptic peptides of KDM4C identified from this analysis is shown in SI Figure S11. The sequence coverage was 91.7% with a redundancy of 1.8%. Mass spectra were processed using the Mass Lynx and DynamX software packages (Waters Corp.). Complete deuteration of control samples was achieved by incubation of 60 pmol KDM4C in the deuterated buffer in the presence of 6 M guanidine for 24 h at 30 °C. Average back exchange (i.e., deuteration loss) was measured as 37% for the analyzed peptides. However, no corrections were made for this deuteration loss as only the relative levels of deuteration incorporation of all samples were compared. The HDX of KDM4C in the presence and absence of peptide after 10 min exchange was measured in triplicate to confirm the significance of the detected changes in deuteration uptake. Significant changes were defined as 3× SD. HDX-MS analysis of KDM4C was also performed in the presence of the nonbinding Peptide 1 and no significant changes in HDX was observed in KDM4C due to the presence of Peptide 1 (data not shown). For peptide 21, KDM4C fragment 331–344 showed complex bimodal HDX kinetics that require further investigation (data not shown). Protein structures were visualized using PyMOL (DeLano Scientific).
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