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Lysine succinylation on non-histone chromosomal protein HMG-17 (HMGN2) regulates nucleosomal DNA accessibility by disrupting the HMGN2–nucleosome association†

Yihang Jing,‡ Gaofei Tian,‡ Xiaoyu Qin, Zheng Liu* and Xiang David Li *

Lysine succinylation (Ksucc) is a novel posttranslational modification that frequently occurs on chromatin proteins including histones and non-histone proteins. Histone Ksucc affects nucleosome dynamics by increasing the DNA unwrapping rate and accessibility. However, very little is known about the regulation and functions of Ksucc located on non-histone chromosomal proteins. Here, we site-specifically installed a succinyl lysine analogue (K_csucc) onto the non-histone chromosomal protein HMG-17 (HMGN2) to mimic the natural succinylated protein. We found that the incorporation of K_csucc into HMGN2 at the K30 site (HMGN2K₃₀succ), which is located within the nucleosome-binding domain (NBD), leads to significantly decreased HMGN2 binding to the mononucleosome. HMGN2K₃₀succ also increased the nucleosomal DNA accessibility by promoting nucleosomal DNA unwrapping in the entry/exit region. This study reveals a novel mechanism of non-histone protein succinylation on altering chromatin recruitment, which can further affect nucleosome and chromatin dynamics.

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Introduction

In eukaryotic cells, genomic DNA is packaged with histones and non-histone proteins into nucleosomes that form chromatin, a highly dynamic structure of repeating nucleosome units. Non-histone chromosomal proteins play vital roles in the maintenance of the chromatin structure and the regulation of chromatin-templated biological processes such as gene expression, DNA replication, and DNA damage repair.

High mobility group nucleosome-binding (HMGN) protein is a family of architectural non-histone chromosomal proteins ubiquitously expressed in vertebrate cell nuclei. HMGs bind specifically to nucleosome core particles with high affinity. The binding of HMGs to nucleosomes is mediated by their evolutionarily conserved and highly positively charged nucleosome binding domain (NBD).¹ By weakening the binding of linker histone H1 to nucleosomes, HMGs destabilise higher-order chromatin structures to promote the binding of nuclear regulatory factors and enhance chromatin-templated transcription activity.^{2,3} Moreover, HMGs can modulate the level of histone PTMs and facilitate the repair of DNA lesions.^{4,5}

Similar to histone, a variety of PTMs have been identified in HMGs including acetylation,^{6,7} phosphorylation,⁸ and SUMOylation,⁹ which regulate their biological activities. These PTMs can often facilitate the dissociation of HMGs from the nucleosome, as they alter the physical and chemical properties of the modified residues. For example, phosphorylation at S24 and S28 of HMGN2, a member of the HMGN family, will induce the dissociation of HMGN2 from the nucleosome due to an unfavourable electrostatic repulsion between NBD and the nucleosomal acidic patch.⁸ Both acetylation and SUMOylation on HMGN2 result in a lower binding affinity to nucleosome core particles, suggesting that PTMs can disrupt the interaction between HMGN2 and the nucleosome.

Besides the aforementioned PTMs, lysine succinylation (Ksucc) has been recently discovered on the K17 and K30 residues of HMGN2, but the biological significance of these non-histone protein PTMs remains unclear.¹⁰ Ksucc frequently occurs on multiple mitochondrial and nuclear proteins and has been shown to have widespread roles in regulating diverse metabolic pathways.^{11,12} Importantly, Ksucc on core histones (e.g., H2BK34 and H4K77) change the lysine charge status from +1 to −1 due to the presence of a carboxylate group, disrupting DNA–histone ionic interactions, leading to reduced nucleosome stability and promoting DNA unwrapping.^{13,14} However, whether and how Ksucc on non-histone chromosomal proteins affects nucleosome dynamics has not yet been studied.

To fill this knowledge gap, we focused on the succinylation of the K17 and K30 sites of HMGN2 to investigate their effects

Department of Chemistry, The University of Hong Kong, Pokfulam Road, Hong Kong, China. E-mail: lz0418@hku.hk, xiangli@hku.hk

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‡ Y. J. and G. T. contributed equally to this work.

on HMGN2 binding to nucleosomes and their potential roles in regulating nucleosome dynamics. Specifically, we employed a well-developed “thiol–ene” reaction to generate succinylated HMGN2 mimics by incorporating a Ksucc analogue (K_{succ}) at K17 and K30 sites to yield HMGN2 $K_{17}\text{succ}$ and HMGN2 $K_{30}\text{succ}$, respectively. We performed biochemical and biophysical characterizations of the interactions between succinylated HMGN2 mimics and nucleosomes, which showed that the K30 succinylation disrupted the binding of HMGN2 to the nucleosome, resulting in increased nucleosomal DNA accessibility and exposing the DNA to chromatin binding proteins such as transcription factors.

Experimental

General methods

All the buffers were prepared with ddH_2O using the standard methods. The synthesized proteins were purified on a preparative high performance liquid chromatography (HPLC) system with a Waters 2535 Quaternary Gradient Module, a Waters 515 HPLC pump, a Waters SFO system Fluidics Organizer and a Waters 2767 Sample Manager. The synthesis of compound **1** was performed following a previously reported method.¹⁴

Preparation of *Xenopus laevis* histones and HMGN2 and LexA proteins

Mutations on K119C in H2A and on K17 or K30 in HMGN2 were introduced by site-directed mutagenesis. All recombinant histones (H2A, H2B, H3, H4, and H2A K119C) and HMGN2 proteins were expressed and purified as previously described.^{15,16} The LexA protein was expressed and purified from the pJWL288 plasmid (kindly gifted by Prof. Michael G. Poirier, Department of Physics, Department of Chemistry & Biochemistry, Ohio State University) as previously described.¹⁷

Preparation of succinylated HMGN2 mimics

Mutated HMGN2 (HMGN2K17C, HMGN2K30C, and HMGN2K17/30C) were dissolved in 0.1 M phosphate buffer (pH 7.0) containing 6 M guanidinium hydrochloride (GdnHCl) to a final concentration of 1 mM. Dithiothreitol (DTT) was added to the reaction mixture to a final concentration of 5 mM, followed by 100 equivalents of compound **1**. Finally, 2,2'-azobis[2-(2-imidazolin-2-yl)propane] dihydrochloride (VA-044) and dimethyl sulphide were added to the reaction mixture at the final concentrations of 20 mM and 100 mM, respectively. The reaction was allowed to take place in a 42 °C water bath for 2 h. The reaction products were analyzed by LC-MS using a Vydac 218 TP C4 column (4.6 mm × 250 mm, Grace) and ESI-MS. The *t*Bu-protected succinylated HMGN2 proteins were purified by preparative reverse phase HPLC (22 mm × 250 mm, Vydac C4, Grace). The pure fractions were pooled and freeze-dried. The lyophilized *t*Bu-protected succinylated HMGN2 proteins were dissolved in 0.1 M phosphate buffer (pH 7.0) containing 6 M GdnHCl to a final concentration of 5 mM. In the deprotection step, 70% trifluoroacetic acid (TFA) was added dropwise to the mixture in an ice-cold bath and left for 1 h at

room temperature with stirring. The deprotected succinylated HMGN2 proteins were then purified by preparative reverse phase HPLC (Vydac C4 column, Grace). The purity and identity of final succinylated HMGN2 mimics were confirmed by ESI-MS (Fig. 2).

Preparation of histone H2A–H2B dimers and (H3–H4)₂ tetramers

The histone dimers and tetramers were prepared following a previously reported method.¹⁵ Briefly, for H2A–H2B dimers, equal amounts of H2A and H2B were mixed and dissolved in unfolding buffer (6 M GdnHCl, 10 mM Tris–HCl, pH 7.5, 1.0 mM EDTA, and 10 mM DTT) to a final histone concentration of 1 mg mL^{−1}. The mixture was incubated on ice for 30 min, followed by dialysis in a histone refolding buffer (2 M NaCl, 10 mM Tris–HCl, pH 7.5, 1.0 mM EDTA, and 5 mM 2-mercaptoethanol) at 4 °C. The refolded dimers were purified using a Superdex 200 pg column (HiLoad 16/60, GE). The same method was used to prepare (H3–H4)₂ tetramers and Cy5-labelled H2A–H2B dimers (H2AK119Cy5–H2B).

Isothermal titration calorimetry (ITC) experiments

All ITC experiments were performed at room temperature (25 °C) in a buffer (10 mM KCl, 10 mM Tris, pH 7.5, and 0.1 mM EDTA) using a MicroCal PEAQ-ITC. The reaction cell containing 200 μL of 15 μM purified nucleosomes was titrated with the corresponding HMGN2 proteins (450 μM). The titration was started with an initial injection volume of 0.4 μL over 0.8 s, followed by 18 injections (2.0 μL) over 4.0 s spaced at intervals of 150 s for each injection. The reference power was set at 10.0 μcal s^{−1} and the stirring speed was 750 rpm. The binding isotherm fit was performed using MicroCal PEAQ-ITC analysis software with a single set of sites to determine the thermodynamic binding constants and stoichiometry.

Electrophoretic mobility shift assay (EMSA) to examine the HMGN2–nucleosome association

The DNA and histone dimers and tetramers were assembled into nucleosomes by step-wise dialysis as previously reported.¹⁸ Briefly, purified histone H2A–H2B dimers and tetramers were incubated with 153 bp of SELEX-generated “Widom 601” DNA (molar ratio of dimer:tetramer:DNA = 2:1:1) at a concentration of 2 μM in 50 μL of reconstitution buffer (2 M KCl, 10 mM Tris, pH 7.8, 0.1 mM EDTA). The samples were transferred into a Slide-A-Lyzer MINI dialysis unit (Thermo) and dialyzed at 4 °C against the reconstitution buffer (concentrations: 1.6 M KCl, 1.4 M KCl, 1.2 M KCl, 1.0 M KCl, 0.8 M KCl, 0.5 M KCl, and 0.2 M KCl) with 50 min for each buffer, followed by 10 mM KCl overnight. The reconstituted nucleosomes were incubated with different amounts of the various HMGN2 proteins in 0.5× TE buffer containing 75 mM NaCl for 15 min at room temperature. The HMGN2–nucleosome mixture was resolved by native-PAGE (5% TBE gel, acrylamide:bis-acrylamide = 29:1). The gel was stained with ethidium bromide (EB) for 15 min and the nucleosome or HMGN2–nucleosome complexes were visualized under UV light using a MyECL Imaging system (Thermo Fisher Scientific).



Preparation of fluorophore-labelled nucleosomal DNA

The fluorophore-labelled DNA fragments were prepared by PCR using primers labelled with Cy3 as previously reported.^{14,19–21} The template for the Cy3-labelled DNA used “Widom 601” DNA with base pairs 8–27 replaced with a LexA recognition sequence (TACTGTATGAGCATACAGTA). The PCR products were precipitated with isopropyl alcohol, washed with ethanol dissolved in water, and then purified using a size-exclusion column (GE Superose 6 increase 10/300). The DNA concentration was determined using a UV-vis spectrophotometer (260 nm, NanoDrop 2000, Thermo).

Preparation and purification of fluorophore-labelled nucleosomes

The H2AK119Cy5–H2B dimers and (H3–H4)₂ tetramers were incubated with Cy3-labelled nucleosomal DNA (molar ratio of dimer:tetramer:DNA = 4 : 1 : 1, with excess amounts of dimers to ensure a fully compacted nucleosome is generated) at a concentration of 2 μM in 50 μL of reconstitution buffer. The dialysis procedure was the same as that for the nucleosomes described above. The reconstituted nucleosomes were purified by native-PAGE (5% TBE gel, acrylamide:bis-acrylamide = 60 : 1) using a Mini Prep Cell (Bio-Rad, Model 491). The purity of nucleosomes used for the FRET assay was checked by native-PAGE (Fig. S1a, ESI[†]).

FRET measurement of nucleosomal DNA unwrapping by LexA titration

The accessibility of DNA in the entry/exit region was determined by the reduction in FRET efficiency as LexA binds to its target sites on the nucleosome.^{21–23} In these experiments, 10 nM Cy3–Cy5-labelled nucleosomes were incubated with HMGN2 in 0.5 × TE buffer containing 75 mM NaCl. The LexA protein was titrated into the system and then incubated at room temperature for 10 min. The Cy3–Cy5-labelled nucleosomes were excited using a 531/25 nm bandpass filter for Cy3 and a 635/25 nm bandpass filter for Cy5. The acceptor (Cy5) emission was collected using a 670/25 nm bandpass filter. For each sample, the fluorescence intensity was measured using a plate reader (Victor X5, PerkinElmer) in triplicate. The FRET efficiency was calculated using the (ratio)_A method as previously reported²⁴ using the equation:

$$E = 2[\varepsilon^A(v'')F^A(v')/F^A(v'') - \varepsilon^A(v')]/[\varepsilon^D(v')d^+]$$

The normalized FRET efficiencies for each LexA concentration were fitted to a non-competitive binding curve:

$$E = E_0 + (E_F - E_0)/(1 + S_{1/2}/[LexA]),$$

where E is the normalized FRET efficiency at a concentration of LexA protein, E_0 is the normalized FRET efficiency in the absence of LexA protein, E_F is the normalized FRET efficiency at a high LexA concentration, and $S_{1/2}$ is the LexA concentration at which the normalized FRET efficiency is reduced by half.

Results and discussion

The synthesis of site-specific succinylated HMGN2 mimics

The study of HMGN2 succinylation *in vitro* requires the generation of homogeneous HMGN2 proteins that carry site-specific

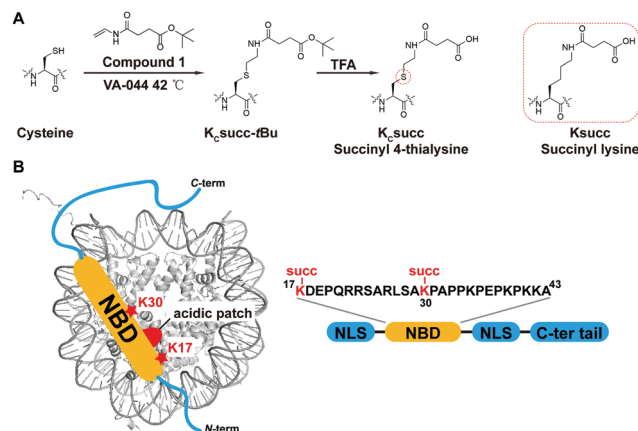


Fig. 1 Illustration of the ‘thiol-ene reaction’ for the installation of lysine succinylation marks. (A) Scheme for the installation of the succinyl lysine analogue (K_c succ) at a cysteine residue. (B) The association of HMGN2 is mediated by its nucleosome-binding domain (NBD). The two reported succinylation sites of HMGN2, K17succ and K30succ, are located within the NBD.

succinylations at the stoichiometric level. In our previous study, we developed a cysteine-selective ‘thiol-ene reaction’ to prepare histone H2B carrying a site-specific lysine succinylation analogue (K_c succ) at K34 (H2BK₃₄succ) (Fig. 1A). Inspired by our successful incorporation of K_c succ onto this histone, we aimed to install K_c succ onto HMGN2, a non-histone protein without cysteine residues. To this end, we focused on two reported succinylation sites on HMGN2, lysine 17 (HMGN2K17) and lysine 30 (HMGN2K30), both located on the nucleosome binding domain (NBD) (Fig. 1B). Therefore, succinylation at these two sites will likely interfere with the interactions between NBD and the nucleosome, resulting in altered nucleosome dynamics.

To introduce succinylation into HMGN2, we first obtained recombinant HMGN2 mutants with the site-specific lysines being replaced by cysteines (*i.e.*, HMGN2K17C, HMGN2K30C, and HMGN2K17/30C double K-to-C mutation). The subsequent thiol-ene reaction on the HMGN2 mutants using compound **1** yielded the *t*Bu-protected succinylated HMGN2 mimic. Deprotection of the *t*Bu-protected product with 70% TFA afforded the desired HMGN2 mimic. The purity and identity of the modified HMGN2K₁₇succ, HMGN2K₃₀succ, and HMGN2K_{17/30}succ proteins were examined and confirmed by LC-MS (Fig. 2).

Succinylation reduces the HMGN2 binding affinity to nucleosomes

We next investigated the effects of HMGN2 succinylation on its interaction with nucleosomes. The non-histone protein HMGN2 normally binds to mononucleosomes to form a complex containing one nucleosome and two molecules of HMGN2.²⁵ The extent of the assembly of the HMGN2–nucleosome complex can be examined by the gel electrophoresis mobility shift assay (EMSA), in which the inverse proportional mobility can reveal differences in the mass of the particles. For this study, unmodified or succinylated recombinant HMGN2 was incubated with *in vitro* reconstituted mononucleosomes in different molar ratios, respectively. As shown



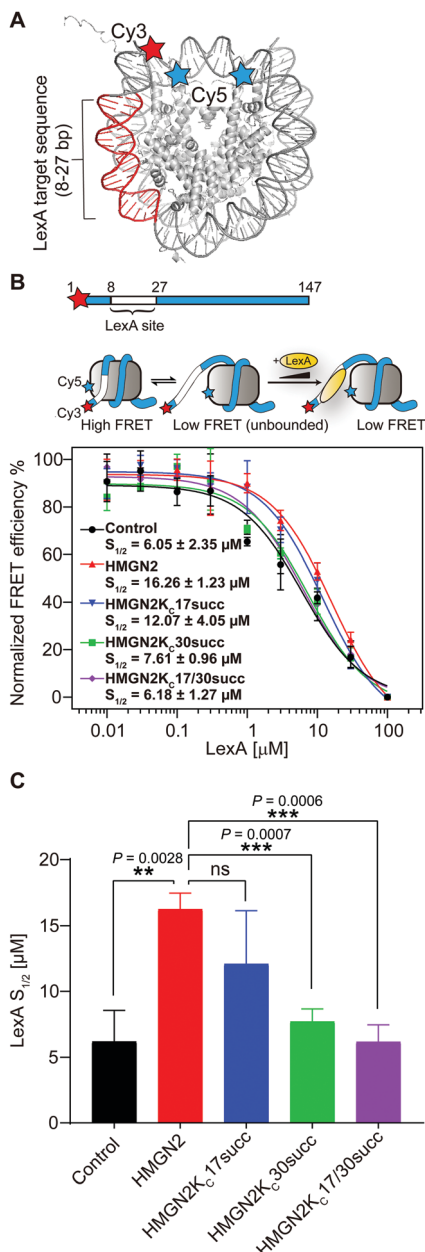


Fig. 5 The succinylation at K30 on HMGN2 increases nucleosomal DNA accessibility. (A) Crystal structure of the nucleosome (PDB: 3LZ1). The locations of the Cy3/Cy5 FRET pair and LexA binding sites. (B) The nucleosomal DNA used for the FRET assay is a 147 bp Widom 601 DNA with a LexA binding site from 8–27 bp and labelled with Cy3 at the 5' end. The three-state model showed decreasing FRET efficiency as the LexA is titrated in and binds to its target site. The plot shows the normalised FRET intensity as a function of LexA concentration for the nucleosome without HMGN2 treatment (Black) or treated with unmodified HMGN2 (red), HMGN2K₁₇succ (blue), HMGN2K₃₀succ (green), and HMGN2K_{17/30}succ (purple). The LexA concentration at which the FRET decreases by 50% is denoted as $S_{1/2}$. For visualization, all curves were normalized between 100% at the maximum FRET value and 0% at the highest LexA concentration. Titrations were repeated in triplicate for each experiment ($n = 3$, $S_{1/2} = \text{mean} \pm \text{SD}$). (C) The histogram shows the $S_{1/2}$ values of nucleosomes ($n = 3$, $S_{1/2} = \text{mean} \pm \text{SD}$). The P values were based on a two-tailed Student's t test. $**P < 0.01$. $***P < 0.001$. ns, no significance.

the $S_{1/2}$ values of HMGN2-treated nucleosomes also increased (Fig. S1B, ESI†). These results verified our hypothesis that the nucleosomal DNA at the entry/exit site in the HMGN2–nucleosome complex is constrained by HMGN2, resulting in decreased DNA accessibility.

We then investigated the effects of HMGN2 succinylation on regulating nucleosomal DNA accessibility using the same experimental design with the three succinylated HMGN2 (*i.e.*, HMGN2K₁₇succ, HMGN2K₃₀succ, and HMGN2K_{17/30}succ). The results showed that the presence of the K_csucc mark at K30 of HMGN2 substantially decreased the nucleosome $S_{1/2}$ value to $7.61 \mu\text{M}$ compared with the unmodified HMGN2, and also approached the level in HMGN2-free nucleosomes ($6.05 \mu\text{M}$). This finding suggests that the HMGN2K₃₀succ shifts the nucleosomal DNA breathing equilibrium toward the unwrapped state, thereby increasing the accessibility of DNA in the entry/exit region. On the other hand, the introduction of succinylation on K17 of HMGN2 (HMGN2K₁₇succ) caused a moderate decrease in the $S_{1/2}$ value of the treated nucleosome compared with the unmodified HMGN2-treated nucleosome. Meanwhile, HMGN2K_{17/30}succ with dual K_csucc marks resulted in only a slight decrease in the $S_{1/2}$ value compared to the HMGN2K₃₀succ-treated nucleosome ($6.18 \mu\text{M}$ vs. $7.61 \mu\text{M}$). These results suggest that lysine succinylation on K30 of HMGN2 plays a predominant role in increasing nucleosomal DNA accessibility.

Conclusions

In this study, we expanded the application of the 'thiol-ene reaction' to prepare site-specific succinylated non-histone proteins (HMGN2K₁₇succ, HMGN2K₃₀succ, and HMGN2K_{17/30}succ). The results of EMSA and ITC assays revealed the introduction of negatively charged lysine succinylation on K30 of HMGN2 changing the net charge of the modified lysine residues from +1 to −1 and significantly disrupting the association of HMGN2 with the mononucleosome. Besides, the FRET-based assay demonstrated that the NBD domain of HMGN2 was required in binding to the acidic patch on the histones and the DNA in the entry/exit region, which decreased the nucleosomal DNA accessibility by constraining the nucleosomal DNA breathing. On the contrary, succinylation on K30 of HMGN2 introduces a negatively charged Ksucc mark, leading to increased nucleosomal DNA accessibility (Fig. 6). Besides K17 and K30 sites on HMGN2, many other lysine succinylation sites have

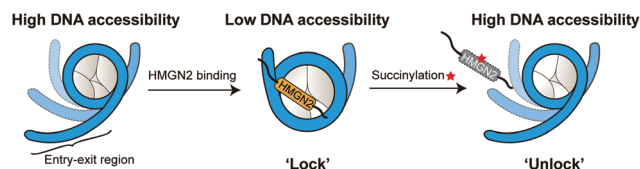


Fig. 6 The function of lysine succinylation in regulating the nucleosomal DNA accessibility by promoting the release of HMGN2 from nucleosomes. In the HMGN2–nucleosome complex (middle), the DNA in the entry/exit region is constrained by HMGN2, which results in a decrease of DNA accessibility. The succinylation on K30 induces the dissociation of HMGN2 from the nucleosome and 'unlocks' the DNA to be accessible.



been identified in several non-histone proteins.^{12,27} A comprehensive investigation of non-histone lysine succinylation will help us to better understand the novel functions of this PTM in regulating chromatin-templated biological processes.

Author contributions

Z. L. and X. D. L. conceived and designed the project. Y. J. synthesized the small-molecule compounds and performed the protein chemistry, nucleosome reconstitution, and FRET experiments. G. T. purified the LexA protein. X. Q. performed the ITC experiments. Z. L. and X. D. L. wrote the manuscript with input from G. T.

Conflicts of interest

There are no conflicts of interest to declare.

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