Deacetylase-independent function of SIRT6 couples GATA4 transcription factor and epigenetic activation against cardiomyocyte apoptosis

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ABSTRACT

SIRT6 deacetylase activity improves stress resistance via gene silencing and genome maintenance. Here, we reveal a deacetylase-independent function of SIRT6, which promotes anti-apoptotic gene expression via the transcription factor GATA4. SIRT6 recruits TIP60 acetyltransferase to acetylate GATA4 at K328/330, thus enhancing its chromatin binding capacity. In turn, GATA4 inhibits the deacetylase activity of SIRT6, thus ensuring the local chromatin accessibility via TIP60-promoted H3K9 acetylation. Significantly, the treatment of doxorubicin (DOX), an anticancer chemotherapeutic, impairs the SIRT6–TIP60–GATA4 trimeric complex, blocking GATA4 acetylation and causing cardiomyocyte apoptosis. While GATA4 hyperacetylation-mimic retains the protective effect against DOX, the hypoacetylation-mimic loses such ability. Thus, the data reveal a novel SIRT6–TIP60–GATA4 axis, which promotes the anti-apoptotic pathway to prevent DOX toxicity. Targeting the trimeric complex constitutes a new strategy to improve the safety of DOX chemotherapy in clinical application.

INTRODUCTION

Anthracyclines, including doxorubicin (DOX), are widely used as efficacious chemotherapeutics, but cause severe dose-dependent cardiotoxicity (1,2). Anthracycline-induced heart failure is largely due to myocyte apoptosis (3). GATA4, a member of the GATA transcription factor family with the ability to bind the consensus DNA motif ‘GATA’, plays critical roles in myocardial differentiation and function (4–6). GATA4 activates the transcription of anti-apoptotic gene Bcl-2 and Bcl-xL etc., which protect against myocyte death induced by DOX (3,7–9). Upon the DOX treatment, GATA4 is rapidly downregulated at both both transcript and protein levels (3,10–12). Intriguingly, GATA4 overexpression causes cardiac hypertrophy (13). These findings suggest that GATA4 might undergo additional layers of fine-tuned regulation, which merits further examination before applying GATA4 restoration as a clinical strategy to prevent DOX-induced cardiotoxicity (3,7,14).

SIRT6 belongs to the highly conserved family of NAD+-dependent sirtuins, which deacetylate histones and non-histone substrates to modulate chromatin stability and restrict transcription (15–17). Through these functions, SIRT6 maintains organismal health and protects against aging and various diseases, including cancers and metabolic disorders (18–21). Cardiac Sirt6 is sensitive to stress stimuli, i.e. angiotensin II, isoproterenol and ischemia/reperfusion-induced reactive oxygen species (ROS) and DOX (23,26–28). Exercise during pregnancy protects neonatal cardiomyocytes against DOX toxicity, accompanied by the increased

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expression of SIRT6 (29). Despite these advances, how SIRT6 protects cardiomyocytes against DOX are unclear.

Here, we demonstrated a novel, deacetylase-independent mechanism by which SIRT6 protects against DOX-induced cardiomyocyte death. Our data suggest that targeting the non-catalytic function of SIRT6 may enhance the safety of DOX chemotherapy.

MATERIALS AND METHODS

Cell culture and treatments

HEK293 (CRL-1573) and H9C2 (GNR-5) cells were purchased from ATCC. Wild-type (WT) and Sirt6+/− mouse embryonic fibroblasts (MEFs) were obtained as previously described (30). Sirt6 knockout (KO) HEK293 cell lines were generated using the CRISPR/Cas9 system, as described previously (21). Primary neonatal mouse cardiomyocytes were prepared with a standard procedure (31). Briefly, hearts from 1- to 3-day-old C57BL/6 mice were isolated and incubated with digestion medium. After centrifuging and plating, the viable cardiomyocytes formed a monolayer with synchronized beating within two days of culture. All cell lines were cultured in Dulbecco’s modified Eagle’s medium (DMEM, Life Technologies, USA) supplemented with 15% fetal bovine serum, 100 U/ml penicillin and streptomycin at 37°C in 5% CO2 and atmospheric oxygen. The cells were treated with DOX at the indicated doses for specific analyses.

Mice and DOX administration

Sirt6+/− mice were crossed with Myh6-cre/Er81 mice to generate Sirt6+/−;Myh6-cre mice. To generate cardiac-specific Sirt6 KO mice, 4-hydroxytamoxifen was injected intraperitoneally (i.p.) daily in Sirt6+/−;Myh6-cre mice for three consecutive days. After 1 month, the heart tissues were harvested and analyzed by PCR-based genotyping and quantitative PCR (qPCR). All mice were housed and handled in the Laboratory Animal Research Center of Shenzhen University. All experiments were performed in accordance with the guidelines of the Institutional Animal Care and Use Committee (IACUC).

Plasmids and RNA interference

Human FLAG-GATA4 and FLAG-TIP60 constructs were purchased from Vigene Biosciences (China). Human FLAG-SIRT1, 2, 6 and 7 constructs were purchased from Addgene (USA). FLAG-GATA4 and FLAG-SIRT6 with amino acid substitutions were generated by PCR-based mutagenesis using a KOD-PLUS kit (Toboyo, Japan). The truncated Sirt6 and GATA4 plasmids were constructed by PCR-based deletion based on the HA-SIRT6 and HA-GATA4 templates. The primers used for mutagenesis are listed in Supplemental Table S1.

For RNA interference, the cells were transfected with small interfering RNAs (siRNAs) for 48 h using Lipofectamine® 3000 (Invivogen, USA), according to the manufacturer’s instructions. Scrambled siRNAs were used as a negative control. All siRNAs were purchased from Gene Pharma (China), and their specific sequences are included in Supplemental Table S1.

Cell transfection and colony formation assay

To establish stably transfected H9C2 cell lines, the pcDNA3.1, GATA4, GATA4 2KR (K328/330R) and GATA4 2KQ (K328/330Q) plasmids were digested with PvuI nuclease for 30 min to linearize, and then individually transfected into H9C2 cells with Lipofectamine® 3000. The medium was replaced after 24 h and supplemented with 2 mg/ml G418 for selection. After 10 days, stably transfected cells were obtained, and their expression was confirmed by western blotting.

For the colony-forming assay, the cells were seeded in six-well plates in triplicate and cultured under normal growth conditions in the presence or absence of DOX at the indicated doses. After culturing for a further 10–14 days, the cell colonies were stained with 0.5% crystal violet solution. The number of colonies in each well was quantitated and the surviving fraction was calculated.

Chromatin-bound fraction assay

The cells were carefully detached from the culture vessel in 1 ml cold PBS buffer and then pelleted by centrifugation at 3000 × g for 1 min. The cell pellets were resuspended with 500 μl Buffer A (10 mM HEPES, 10 mM KCl, 1.5 mM MgCl2, 0.34 M sucrose, 10% glycerol, 1 mM DTT, 0.1% Triton X-100, 10 mM Na3VO4 and protease inhibitor cocktail) and incubated on ice for 10 min. The cell lysates were then centrifuged at 1300 × g for 5 min and supernatant containing the cytosolic proteins was collected. The pellet (chromatin-bound fraction) was obtained and denatured with 1 × SDS loading buffer.

Protein extraction, western blotting and antibodies

The mouse tissues were homogenized in 1 ml ice-cold tissue lysis buffer (25 mM Tris–HCl, pH 7.5, 10 mM Na3VO4, 100 mM NaF, 50 mM Na4P2O7, 5 mM EGTA, 5 mM EDTA, 0.5% SDS, 1% NP-40 and protease inhibitor cocktail). Following sonication, the lysates were centrifuged at 16 000 × g for 10 min and then carefully collected. The protein concentrations were determined using a Bicinchoninic Acid kit (BCA; Thermo). Western blotting was performed following a standard protocol. Anti-acetyl lysine (CST, Ac-K-100, #6952), anti-poly/mono ADP ribose (CST, #83732) and anti-Tip60 (CST, #12058) antibodies were purchased from Cell Signaling Technology (USA). Anti-GATA4 (ab84593), anti-SIRT6 (ab62739) and anti-H3 (ab1791) antibodies were purchased from Abcam (UK). Anti-P300 (sc48343) and anti-GST (sc138) antibodies were purchased from Santa Cruz Biotechnology (USA). Anti-H3K9ac (ABE18) was obtained from EMD Millipore (USA) and anti-FLAG M2 was purchased from Sigma-Aldrich (USA).

Co-immunoprecipitation (Co-IP) and pull-down assays

The co-immunoprecipitation assay was performed as previously described (32). Briefly, the cells were washed with cold

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PBS and harvested. The cell pellets were then resuspended with 500 μl IP lysis buffer (20 mM Tris–HCl PH 8.0, 250 mM NaCl, 5 M sodium pyrophosphate, 1 mM sodium fluoride, 10% glycerol, 2 mM EDTA, 10 mM Na3VO4, protease inhibitor cocktail and 1 mM nicotinamide). Following sonication and centrifugation, the supernatant was collected and incubated with anti-HA or anti-FLAG M2 agarose (Sigma, USA) overnight at 4°C on a rotating platform. The beads were then washed three times with 1 ml cold lysis buffer. To denature, 1× SDS loading buffer was added to the samples and boiled at 98°C for 6 min. The samples, including the input (5%), were then analyzed by western blotting.

For the in vitro pull-down assay, GST or GST-SIRT6 (1 μg)-conjugated GSH beads were suspended in NETN buffer (20 mM Tris–HCl pH 8, 100 mM NaCl, 1 mM EDTA, 0.5% NP40 and phosphatase and protease inhibitor cocktail). The purified His-GATA4 proteins (0.2 μg) were then added and incubated overnight at 4°C, followed by three washes with NETN buffer. The boiled eluates were then separated and detected by Coomassie staining and western blotting.

Luciferase reporter assay
HEK293 cells were transfected with a pGL3-Bcl-2-promoter reporter, together with the expression plasmids and a control pCMXβ-gal plasmid. The luciferase activity was measured using a luciferase reporter assay kit (Promega, #E1501), according to the manufacturer’s instructions. The luciferase values were normalized according to β-gal activity.

RNA preparation and real-time qPCR analysis
Total RNAs were extracted from cells or mouse tissues using Trizol® reagent RNAiso Plus (Takara, Japan) following the phenol-chloroform extraction method. A cDNA library was constructed from the purified RNAs using Prime Script RT Master Mix (Takara, Japan) and the following thermal cycling conditions: 37°C for 40 min; 85°C for 5 s. Gene expression was analyzed using a CFX Connected™ Real-time PCR Detection System (Bio-Rad, USA) with SYBR Ex Taq Premixes (Takara, Japan). Each experiment was performed at least three times. The values were standardized to 18s rRNA. The primers for qPCR are listed in Supplemental Table S1.

Chromatin immunoprecipitation (ChIP) assay
The ChIP assay was performed as previously described (21). Briefly, the cells were cross-linked and lysed with lysis buffer (50 mM Tris–HCl pH 8.0, 2 mM EDTA, 15 mM NaCl, 1% SDS, 0.5% deoxycholate, protease inhibitor cocktail and 1 mM PMSF). Following sonication and centrifugation, the supernatant was collected and precleared with protein A/G Sepharose. The precleared samples were incubated overnight with the indicated antibodies: anti-H3K9ac, anti-SIRT6, anti-GATA4 antibody (2 μg/sample) and appropriate control IgGs. The bound DNA fragments were precipitated by supplementing with 50% suspension of protein A/G Sepharose. Following washing sequentially with a series of buffers, the beads were heated to 65°C to reverse the cross-links. The DNA fragments were purified and analyzed by qPCR. The primers used in this study are listed in Supplemental Table S1.

TUNEL staining assay
DOX-induced cardiomyocyte apoptosis was evaluated using a Dead End™ Fluorometric TUNEL System (Promega, G3250), according to the manufacturer’s instructions. Briefly, the cells were fixed and incubated with a reaction mixture containing terminal reoxynucleotidyl transferase and Fluorescein-12-dUTP at 37°C for 1 h. DAPI was used to counterstain the nuclei. Five fields-of-view were randomly selected, and images were captured under the microscope to calculate the TUNEL-positive staining rate for each slice.

Fluorescence-activated cell sorting (FACS) assay
The H9C2 cells were treated with 1 μM DOX for 12 h and then stained with FITC-Annexin V (Beyotime, C1062L) and 7-AAD (BD Pharmingen) for live/dead cell discrimination. Flow cytometry analyses were performed using an LSR-II flow cytometer and analyzed with FlowJo x7.5.

In vitro SIRT6 deacetylation assay
SIRT6 deacetylation assay was performed with the SIRT6 Activity Assay Kit (Fluorometric) (ab 156068, Abcam, UK) following the manufacturer’s instructions. Briefly, bacterially purified GST-SIRT6 or GST-SIRT6 H133Y was pre-incubated with purified His-GATA4, G4-13 peptide (RKKRPKNLNKSKT) or G4-24 peptide (ACGLYMKL-HGVPRLAMRKEGIQT) (GL Biochem LTD, Shanghai, China), in the reaction buffer at 4°C for 30 min. Once the fluorophore-conjugated synthetic Ac-peptide was added, the plate was moved to measure the fluorescence intensity.

Statistical analyses
All values are expressed as the mean ± s.e.m. Statistical differences among groups were determined using Student’s t-test or two-way ANOVA using GraphPad Prism software. Survival curve data were analyzed using a log-rank (Mantel–Cox) test.

RESULTS
Sirt6 enhances anti-apoptotic gene expression against DOX via Gata4
To determine how Sirt6 protects cell survival against DOX, we assessed the RNA-seq data sets from two independent Sirt6 null cell lines: mouse embryonic stem cells (mESCs) (33) and mouse embryonic fibroblast cells (MEFs) (21). As shown, Sirt6 depletion activates apoptotic pathway, with dramatic decrease of anti-apoptotic gene expression, like Bcl-2 and Bcl-xL (Supplemental Figure S1A, B). GATA4 transcription factor is pivotal for postnatal cardiomyocyte...
survival against DOX by direct targeting the promoter regions of anti-apoptotic genes (3,7–9). Indeed, mRNA levels of \textit{Gata4}, \textit{Bcl}-2 and \textit{Bcl}-xL were all rapidly downregulated in H9C2 cells and primary mouse cardiomyocytes upon DOX treatment (Figure 1A and Supplemental Figure S1C). To further understand the causal link between Sirt6 and the \textit{Gata4}/\textit{Bcl}-2 pathway, we generated cardiac-specific \textit{Sirt6} knockout in adult mice using the tamoxifen-inducible α-myosin heavy chain (Myh6) Cre line, i.e. \textit{Sirt6}\textsuperscript{+/−}/\textit{Myh6-Cre}, which spontaneously develop heart failure (23). qPCR analysis showed that \textit{Sirt6} depletion in mouse hearts significantly reduced \textit{Bcl}-2 and \textit{Bcl}-xL levels, with little effect on \textit{Gata4} mRNA or protein expression (Figure 1B and Supplemental Figure S1D). \textit{Sirt6} knockdown by siRNAs further exacerbated the DOX-induced reduction of \textit{Bcl}-2 and \textit{Bcl}-xL mRNA expression in H9C2 cells (Figure 1C and Supplemental Figure S1E). By contrast, \textit{Sirt6} overexpression enhanced \textit{Bcl}-2 and \textit{Bcl}-xL expression levels and protected against the decline induced by DOX in mice (Figure 1D and Supplemental Figure S1F).

We then asked whether \textit{Sirt6} regulates anti-apoptotic pathway via \textit{Gata4}. To this end, we first compared the down-regulated genes in \textit{Sirt6} null mESCs and the genes that are bound by \textit{Gata4} obtained from chromatin immunoprecipitation and sequencing (ChiP-seq) in mESCs-derived mesoderm (34). Intriguingly, almost 50% of down-regulated genes caused by \textit{Sirt6} deficiency harbor \textit{Gata4} binding sites and gene ontology analysis revealed significant enrichment of pathways regulating apoptosis (Supplemental Figure S1G, H and Supplemental Table S2). We thus speculate that \textit{Sirt6} and \textit{Gata4} might cooperate in regulating anti-apoptotic gene expression. As described, \textit{Gata4} regulates cardiac \textit{Bcl-2} transcription by binding to the proximal −266 \textit{Gata4} motif (P1) rather than the distal −1025 \textit{Gata4} motif (P2) (8). Interestingly, ChIP analysis demonstrated that \textit{Sirt6} was also highly enriched at the P1 site (Figure 1E, F). We then determined the luciferase activity under transcriptional control of \textit{Bcl}-2 promoter containing the P1 motif. We found that ectopically expressed \textit{Sirt6} significantly elevated the \textit{Bcl}-2 promoter (P1) -driven luciferase activity (Figure 1G). Moreover, \textit{Sirt6} overexpression enhanced the \textit{Gata4} occupancy at the P1 site (Figure 1H and Supplemental Figure S1I), but \textit{Sirt6} KD compromised it (Figure 1I and Supplemental Figure S1J). Together, these data suggest that \textit{Sirt6} enhances \textit{Gata4} chromatin binding to promote anti-apoptotic gene expression.

**GATA4 physically associates with SIRT6**

As shown above, \textit{Sirt6} and \textit{Gata4} are both enriched at the P1 promoter region and coordinate the transcription of \textit{Bcl}-2, suggesting that these two proteins may directly interact. Indeed, we detected FLAG-GATA4 in anti-\textit{HA-Sirt6} immunoprecipitates in HEK293 cells expressing FLAG-GATA4 together with or without HA-\textit{Sirt6} (Figure 2A). Reciprocally, HA-\textit{Sirt6} was present in the purified pool of FLAG-GATA4 from HEK293 cells expressing HA-\textit{Sirt6} together with or without FLAG-GATA4 (Figure 2B). To confirm such interaction under physiological condition, we did endogenous co-immunoprecipitation (CoIP) and immunofluorescence microscopy in H9C2 cardiomyocytes.

Again, \textit{Sirt6} was detected in the anti-GATA4 immunoprecipitates and co-stained with GATA in the nucleus (Figure 2C, D). Next, we performed \textit{in vitro} pull-down assay using purified \textit{E. coli}-expressing His-GATA4 and GST-SIRT6. As shown, the result suggested a direct interaction between \textit{GATA4} and \textit{Sirt6} (Figure 2E). \textit{GATA4} possesses two zinc finger (Zn-finger) domains; the C-terminal Zn-finger is essential for DNA and cofactor binding (35,36) (Figure 2F). Domain mapping showed that \textit{Sirt6} specifically bound \textit{GATA4} at C-terminal Zn-finger region (a.a. 251–350) (Figure 2G).

**SIRT6 facilitates GATA4 acetylation independently of its deacetylation activity**

We next sought to figure out how \textit{Sirt6} entrains \textit{Gata4}-targeted gene expression. Of note, post-translational modifications, particularly phosphorylation and acetylation, are critical to \textit{Gata4} DNA-binding capacity and activation (37–39). To examine whether \textit{Sirt6} modulates the acetylation level of \textit{Gata4}, we IPed FLAG-GATA4 in HEK293 cells co-overexpressing \textit{GATA4} and \textit{Sirt6} and examined its acetylation level using pan anti-acetyl lysine antibodies. Rather surprisingly, the acetylation level of \textit{Gata4} was markedly increased in the presence of ectopic \textit{Sirt6} (Figure 3A). Low level of \textit{Gata4} acetylation was detected in the \textit{Sirt6}-depleted cells (Supplemental Figure S2A). In addition to deacetylation activity, \textit{Sirt6} also possesses mono-ADP-ribosylase (MAR) activity and could auto-MAR itself (40). However, no \textit{Gata4} MAR were observed, indicating that \textit{Gata4} is less likely a MAR target of \textit{Sirt6} (Supplemental Figure S2B).

We next examined whether the increase of \textit{Gata4} acetylation requires deacetylation activity of \textit{Sirt6}. Nicotinamide (NAM) binds to a conserved pocket adjacent to that of NAD\textsuperscript{+} to block deacetylase and mono-ADP-ribosylase activities of \textit{Sirt6} (41–43). We observed that NAM abrogated sirtuin-mediated H3K9 deacetylation but was unable to inhibit the increased level of \textit{Gata4} acetylation induced by \textit{Sirt6} overexpression (Figure 3B). Further, the catalytically inactive \textit{Sirt6} mutants, D63Y and D116N (26), remained capable of binding to and enhancing \textit{Gata4} acetylation (Figure 3C). More importantly, the overexpression of \textit{Sirt6} WT, D63Y and D116N restored \textit{Bcl}-2 and \textit{Bcl}-xL expression in \textit{Sirt6}-deficient MEFs (Figure 3D and Supplemental Figure S2C), and significantly inhibited apoptosis in H9C2 cells exposed to DOX (Figure 3E, F). These data thus implicate that \textit{Sirt6} promotes \textit{Gata4} acetylation and thus transcription activity independent of its deacetylation activity.

**TIP60 is required for SIRT6-induced GATA4 acetylation**

The data that \textit{Sirt6} interacts with \textit{Gata4} and enhances its acetylation level suggests that additional factors, most likely histone acetyltransferase (HAT), are involved. To determine which HAT is responsible for enhanced \textit{Gata4} acetylation, we examined the effects of \textit{Sirt6}-regulated \textit{Gata4} acetylation upon exposure to a series of HAT inhibitors, including Cpth2 to inhibit Gcn5 (44), Mg149 to inhibit TIP60 (45) and C646 to inhibit p300 (46,47). In-
Figure 1. SIRT6 antagonizes DOX-induced apoptotic pathway in cardiomyocytes via GATA4. (A) Quantitative PCR (qPCR) showing the mRNA levels of Bcl-2, Bcl-xL, Gata4, and Sirt6 in primary mouse cardiomyocytes after the treatment of DOX (1 μM) for 6 h. ***P < 0.001, **P < 0.01, ‘ns’ indicates no significance. (B) qPCR results showing the mRNA levels of Bcl-2, Bcl-xL, Sirt6 and Gata4 in heart tissues isolated from Sirt6+/− and Sirt6−/−;Myh6-Cre mice treated with tamoxifen. *P < 0.05, **P < 0.01, ***P < 0.001, ‘ns’ indicates no significance. (C) H9C2 cells with si-NC (negative control) or si-Sirt6 small interfering (si)RNAs were exposed to DOX (1 μM, 6 h). The mRNA levels of Bcl-2, Bcl-xL and Gata4 were determined by quantitative PCR. *P < 0.05, **P < 0.01, ‘ns’ indicates no significance. (D) qPCR results showing the mRNA levels of Bcl-2, Bcl-xL and Sirt6 in wild-type (WT) and Sirt6 transgenic (Sirt6-TG) mice after intraperitoneal injection of DOX (n = 5/group). *P < 0.05, **P < 0.01, ***P < 0.001, ‘ns’ indicates no significance. (E) Schematic illustration of the GATA consensus motif on Bcl-2 promoter region. P1 and P2 indicate –266 GATA motif and –1025 GATA motif respectively. TSS represents the transcription start site of the Bcl-2 gene. (F) ChIP was performed using anti-SIRT6 antibodies and IgG. Enrichment of SIRT6 on the Bcl-2 promoter region was analyzed by qPCR with primers targeting the P1 and P2 motifs. **P < 0.01. (G) HEK293 cells were transfected with indicated plasmids, and the luciferase activities were measured and normalized to β-gal activities. **P < 0.01. The western blots indicate the expression levels of FLAG-GATA4 and FLAG-SIRT6. (H) ChIP was performed to test GATA4 enrichment on the Bcl-2 promoter using anti-GATA4 antibody in HEK293 cells with or without FLAG-SIRT6 overexpression. *P < 0.05. (I) ChIP was performed to test GATA4 enrichment on the Bcl-2 promoter using anti-GATA4 antibody in HEK293 cells with si-NC or si-SIRT6. *P < 0.05.

Interestingly, while H3K56ac levels were further reduced by the treatment of all HAT inhibitors, only Mg149 treatment significantly inhibited the increased acetylation of GATA4 induced by SIRT6 (Figure 4A). Specific Tip60 deletion in mouse hearts shortens lifespan in mice, accompanied with increased myocyte density and apoptosis (48). Depletion of Tip60–p400 HAT complex severely affected Gata4-regulated primitive endoderm differentiation in ESCs (49). However, whether GATA4 is a direct target of TIP60 remains unknown. We found that TIP60 overexpression increased GATA4 acetylation level, and that this effect was eliminated upon application of Mg149 (Supplemental Figure S3A, B). TIP60 KD by siRNA also markedly decreased GATA4 acetylation levels (Supplemental Figure
Figure 2. SIRT6 physically interacts with GATA4. (A) Co-immunoprecipitation using anti-HA agarose was performed in HEK293 cells transfected with indicated constructs. The level of FLAG-GATA4 was detected by western blotting in purified pool of HA-SIRT6. (B) Co-immunoprecipitation using anti-FLAG agarose was performed in HEK293 cells transfected with indicated constructs. The level of HA-SIRT6 was detected by western blotting in purified pool of FLAG-GATA4. (C, D) Endogenous interaction of GATA4 and SIRT6 in H9C2 cells was evaluated by co-immunoprecipitation (Co-IP) and immunostaining with anti-GATA4 and anti-SIRT6 antibodies. Scale bar, 50 μm. (E) GST pull-down assay was performed to test the invitro binding of purified His-GATA4 and GST-fused SIRT6 from E.coli. Coomassie blue staining indicates the loading levels of GST and GST-SIRT6. (F) Diagram of full-length (F-L) and fragmented human GATA4 with one or two zinc finger DNA-binding domains. (G) Domain-based truncationsof HA-GATA4 were co-expressed with FLAG-SIRT6 in HEK293 cells. HA-GATA4 protein was precipitated and FLAG-SIRT6 was analyzed by western blotting. S3C). These results indicate that TIP60 is a novel acetyltransferase of GATA4.

To confirm the role of TIP60 in SIRT6-enhanced GATA4 acetylation, we examined the acetylation level of GATA4 in TIP60 KD cells with SIRT6 overexpression. We found that the SIRT6-enhanced acetylation of GATA4 was significantly abolished in TIP60 KD cells (Supplemental Figure S3D). In the other way around, overexpression of TIP60 in SIRT6-depleted cells was unable to fully acetylate GATA4 (Figure 4B). Importantly, endogenous Sirt6 and Tip60 were present in the anti-Gata4 immunoprecipitates from primary mouse cardiomyocytes (Figure 4C), implying that these proteins form a trimeric complex. Moreover, low level of GATA4 acetylation in SIRT6 KO cells was associated with an undetectable interaction between TIP60 and GATA4 (Figure 4D). These data suggest that SIRT6 acts as a scaffold to facilitate TIP60-mediated GATA4 acetylation.

**TIP60 acetylates GATA4 at K328/330**

The C-terminal Zn-finger domain of GATA4 is often acetylated (36), which is, as shown above, the binding site for SIRT6. Deletion of the C-terminal Zn-finger domain eliminated both acetylation and chromatin association of GATA4, as determined by anti-AcK and anti-histone H3 immunoblotting in the anti GATA4 immunoprecipitates and the chromatin fractionation assay (Figure 4E and Supplemental Figure S3E). There are six evolutionally conserved lysine (K) residues within this region (Figure 4F). It has been reported that p300 acetylates GATA4 on K313/320/322/324 and thus enhances the DNA-binding activity, resulting in stress-induced cardiac hypertrophy (36,50). However, by K-to-R mutation, we noticed that K313/320/322 were less likely acetylated by TIP60. Instead, the K328/330R mutant (2KR) almost totally abolished the increased GATA4 acetylation induced by either ectopic SIRT6 or TIP60 (Figure 4G and Supplemental Figure S3F). Together, these data suggest that TIP60 mediates GATA4 acetylation at K328/330.

Auto-acetylation activates TIP60, which is negatively regulated by SIRT1 (51). We thus examined whether SIRT6 modulates the acetylation status of TIP60. As shown, SIRT6 overexpression had marginal effect on the acetylation level of TIP60 (Figure 4H). Further Co-IP analy-
Figure 3. SIRT6 enhances GATA4 acetylation and protects against DOX-induced myocyte apoptosis independently of its deacylase activity. (A) Western blotting showing the acetylation level of GATA4 immunoprecipitated from HEK293 cells overexpressing FLAG-SIRT6 and HA-GATA4. Acetylation level was determined with anti-acetyl lysine (AcK) antibodies. Lower, fold change of the acetylation level of GATA4 relative to HA-GATA4 only control, determined by ImageJ. \( n = 6 \). *** \( P < 0.001 \). (B) HEK293 cells co-expressing FLAG-SIRT6 and HA-GATA4 were treated with 10 μM nicotinamide (NAM) for 4 h. The acetylation level of GATA4 was examined using anti-AcK antibodies following immunoprecipitation. Lower, fold change of the acetylation level of GATA4 relative to HA-GATA4 only control, determined by ImageJ. \( n = 3 \). *** \( P < 0.001 \). ‘ns’ indicates no significance. (C) HA-GATA4 co-expressed with different forms of FLAG-SIRT6 (WT, D63Y, D116N and H133Y mutant) separately in HEK293 cells was immunoprecipitated with anti-HA beads. The blot of FLAG-SIRT6 in IPs indicate the association of GATA4 and SIRT6. Lower, fold change of the acetylation level of GATA4 relative to HA-GATA4 only control, determined by ImageJ. \( n = 3 \). ** \( P < 0.01 \); * \( P < 0.05 \). ‘ns’ indicates no significance. (D) Real-time PCR analysis of mRNA levels of Bcl-2 in WT and Sirt6 \(^{-/-}\) MEFs re-expressing FLAG-SIRT6, or enzyme-dead mutations. *** \( P < 0.001 \); *** \( P < 0.001 \). 18srRNA was used as a reference gene. (E, F) Representative flow plots and quantification showing the percentages of apoptotic H9C2 cells overexpressing the indicated genes with the treatment of DOX (1 μM, 12 h) following fluorescence-activated cell sorting assay. * \( P < 0.05 \); ** \( P < 0.01 \); *** \( P < 0.001 \).
Figure 4. SIRT6 recruits TIP60 for GATA4 acetylation at Lysine 328/330. (A) Immunoblot of anti-HA immunoprecipitates showing the acetylation level of HA-GATA4 in HEK293 cells overexpressing ectopic HA-GATA4 treated with indicated histone acetyltransferase (HAT) inhibitors, i.e. Cph2 (20 μM), Mg149 (20 μM) and C646 (10 μM), for 6 h. Lower, fold change of the acetylation level of HA-GATA4 relative to HA-GATA4 only control, determined by ImageJ. n = 3. *** P < 0.001. ‘ns’ indicates no significance. (B) Immunoblot of anti-HA immunoprecipitates showing the acetylation level of HA-GATA4 in HEK293 cells overexpressing ectopic HA-GATA4 together with FLAG-TIP60 or empty vector HA-GATA4 proteins were purified from si-N C- or si-SIRT6-treated HEK293 cells co-expressing HA-GATA4. The acetylation status of GATA4 was determined by western blotting with anti-AcK antibodies. Right, fold change of the acetylation level of HA-GATA4 relative to HA-GATA4 only control, determined by ImageJ. n = 4. *** P < 0.001. (C) Western blotting showing the levels of endogenous Tip60 and Sirt6 in anti-Gata4 IPs from primary mouse cardiomyocytes. IgG serves as an antibody control. (D) FLAG-TIP60 was co-expressed with or without HA-GATA4 in wild-type (WT) or SIRT6 knockout (KO) HEK293 cells. Co-IP analysis of HA-GATA4 was performed and the levels of FLAG-TIP60 and GATA4 acetylation were assessed. Right, fold change of FLAG-TIP60 protein bound to HA-GATA4 relative to WT HEK293 cell control, determined by ImageJ. n = 4. ** P < 0.01. (E) Truncated HA-GATA4 constructs were co-transfected with or without FLAG-SIRT6. Western blotting was performed with the indicated antibodies. Lower, fold change of HA-GATA4 acetylation level relative to HA-GATA4 (F-L) only control, determined by ImageJ. n = 3. *** P < 0.001. ‘ns’ indicates no significance. (F) Protein sequence alignment of C-terminal Zn-finger domain of human GATA4 and its orthologues in mouse, rat and Xenopus. All K residues within this region targeted for acetylation were highlighted. (G) The constructs of WT and site-mutated (K328/330R) HA-GATA4 were co-expressing with FLAG-TIP60 in HEK293 cells. The GATA4 acetylation levels were determined. Right, fold change of HA-GATA4 acetylation level relative to HA-GATA4 WT only control, determined by ImageJ. n = 4. *** P < 0.001. ‘ns’ indicates no significance. (H) Western blotting showing the levels of HA-SIRT6 protein and TIP60 acetylation in HEK293 cells co-transfected with FLAG-TIP60 and HA-SIRT6 or empty vector. Right, fold change of FLAG-TIP60 acetylation level relative to FLAG-TIP60 only control, determined by ImageJ. n = 3. ‘ns’ indicates no significance.
sis showed that both WT and enzyme-dead SIRT6 bound to TIP60 (Supplemental Figure S3G). We also examined the MAR levels of TIP60 when SIRT6 was overexpressed, but hardly observed TIP60 MAR, suggesting that TIP60 is less likely MARed (Supplemental Figure S3H). The data support a deacetylase-independent and structural role of SIRT6 in GATA4 acetylation.

GATA4 acetylation at K328/330 is critical for DOX resistance

Restoring GATA4 transcriptional activity has been highly recognized as a promising approach to prevent DOX-induced heart failure (7,12). We next investigated the car-dioprotective role of TIP60-mediated GATA4 acetylation. We co-expressed HA-GATA4 and FLAG-SIRT6, and then treated the cells with DOX. Significantly, the DOX treatment disrupted the interaction of GATA4 and SIRT6, and simultaneously reduced the levels of GATA4 acetylation in a dose-dependent manner (Figure 5A). In primary cardiomyocyte, DOX treatment dramatically reduced the Bel-2 expression, which was largely rescued by overexpression of GATA4 WT and K328/330Q (2KQ), but not 2KR (Figure 5B). Consequently, overexpression of GATA4 WT or 2KQ was able to alleviate DOX-induced myocyte apoptosis as determined by TUNEL assay, but 2KR mutant failed (Figure 5C). We further generated H9C2 cell lines stably overexpressing GATA4 WT, 2KR mutant or 2KQ mutant (Supplemental Figure S4A). While the ectopic GATA4 WT and 2KQ mutant significantly attenuated the decline of Bel-2 and Bel-xL levels in H9C2 cells exposed to DOX, the 2KR mutant had little effect (Figure 5D and Supplemental Figure S4B). Consistently, ChIP data showed that the 2KR, but not the 2KQ mutant, compromised the DNA-binding capacity of GATA4, either in presence or absence of DOX (Figure 5E). GATA4 overexpression alleviates DOX-induced cardiomyocyte apoptosis and resultant myocardial atrophy; however, this benefit was abrogated by the 2KR mutation (Figure 5F, G and Supplemental Figure S4C, D). Further colony formation assay demonstrated that GATA4 2KQ mutant promoted myocyte survival to a similar extent as WT in response to DOX, whereas the 2KR mutant failed to do so (Figure 5H). These data support the notion that TIP60-mediated acetylation is critical for the cardioprotective role of GATA4.

The SIRT6–TIP60–GATA4 axis epigenetically ensures gene expression

SIRT6 and TIP60 are the main HDAC and HAT for H3K9 and/or H3K56 acetylation, which dictates open chromatin structure to facilitate transcription (52,53). We therefore asked whether/how the SIRT6–TIP60–GATA4 axis integrate chromatin remodeling to gene transcription. Again, using the promoter of Bel2 as an example, the ChIP data showed that TIP60 was highly enriched on the P1 region, which was significantly dampened when SIRT6 was knocked down (Figure 6A). Consistently, overexpression of TIP60 significantly enhanced the level of H3K9ac at P1 (Figure 6B), which was diminished by SIRT6 deple-tion (Figure 6C). Interestingly, despite the global levels of H3K9ac and H3K56ac were remarkably decreased upon ectopically expressed SIRT6 (Supplemental Figure S5A), ChIP assay showed that the enrichment of H3K9ac and H3K56ac at the P1 motif was unchanged (Figure 6C). Likewise, SIRT6 knockdown (KD) unaffected the enrichment of H3K9ac at the P1 motif (Supplemental Figure S5B). By contrast, GATA4 KD significantly downregulated the level of H3K9ac at the promoter locus without affecting SIRT6 and TIP60 enrichment (Figure 6D–F).

We noticed that the enzyme-dead mutant H133Y completely abolished the interaction between SIRT6 and GATA4 (Figure 6G), and the ability against DOX-induced cardiomyocyte apoptosis (Supplemental Figure S5C). We therefore reasoned that the association of GATA4 might suppress deacetylase activity of SIRT6 on the target pro-moter. To gain more experimental evidences, we performed in vitro deacetylation assay and found that purified His-tagged SIRT6 significantly blocked deacetylase activity of recombinant SIRT6 (Figure 6H). Considering that the SIRT6/TIP60 complex mediates GATA4 acetylation on K328/330 and the C-terminal Zn-finger of GATA4 is essential for SIRT6 binding, we synthesized two peptides in this region (a.a. 295–331) of GATA4: ACGLYMKL-HGVPRPLAMRKEGIQT (a.a. 295–318) termed G4-24 and RRKRPKLNKSKT (a.a. 319–331) termed G4-13. The in vitro deacetylation assay showed that the G4-13, but not G4-24, dramatically inhibited the deacetylase activity of SIRT6 (14 μM for 50% inhibition) (Figure 6I and Supple-mental Figure SSD).

Considering that H133 site is also required for the nucleosome-association of SIRT6 (54) and together with our findings, we proposed that SIRT6 occupies the promo-tor region via a general nucleosome-binding property: SIRT6 recruits TIP60, globally or at specific gene promoter, to balance dynamic acetylation of histones; when GATA4 is recruited and blocks SIRT6 deacetylation activity, TIP60 acetylates GATA4 to enhance its transcription activity, and to acetylate histones, e.g. H3K9ac, thus ensuring an open chromatin structure for transcription (Figure 7).

DISCUSSION

The clinical use of DOX against malignancies is limited due to severe adverse effect—cardiotoxicity. Multiple mecha-nisms are involved in the DOX-induced heart failure, including oxidative stress, DNA damage and apoptosis (55,56). In response to DOX, GATA4 is downregulated and fails to promote cardiomyocyte survival (3,7). Here, we revealed that dynamic GATA4 acetylation, which is independent of p300 (36.50), is required for its transac-tivation against DOX-induced cardiotoxicity. SIRT6 acts as a scaffold to recruit TIP60, which acetylates GATA4 to enhance transcriptional activity. SIRT6/TIP60-regulated GATA4 acetylation might be considered as potential targets to prevent DOX cardiotoxicity.

The regional epigenetic regulation is a fundamental mechanism that modulates gene transcription. SIRT6 is well-recognized as a transcriptional repressor, which catalyzes histone deacetylation using NAD+ as a coenzyme or substrate (57,58). The D63Y and D116N mutations disrupt the affinity of SIRT6 for NAD+ (59), thus eliminat-
Figure 5. Hypoacetylation of GATA4 diminishes cardio-protection against DOX. (A) Western blotting analysis of the levels of FLAG-SIRT6 protein and HA-GATA4 acetylation in the precipitated pool of HA-GATA4 with DOX treatment at 0, 1, 2 μM for 6 h (left). Percent level of acetylated GATA4 relative to vehicle/no FLAG-SIRT6 control (middle) and percent FLAG-SIRT6 bound to HA-GATA4 relative to vehicle control (right). Quantification was performed by ImageJ. \( n = 3 \), \( * P < 0.05 \), \( ** P < 0.01 \). (B) qPCR analysis of Bcl-2 mRNA levels in primary mouse neonatal cardiomyocytes ectopically expressing GATA4 WT, K328/330R (2KR) and K328/330Q (2KQ) upon exposure to DOX (1 μM, 6 h). *** \( P < 0.001 \), 'ns' represents no significance. (C) Percent TUNEL positively stained primary cardiomyocytes after treatment with DOX (1 μM) for 24 h. ** \( P < 0.01 \), *** \( P < 0.001 \), 'ns' indicates no significance. (D) H9C2 cells stably expressing GATA4 WT, K328/330R (2KR) and K328/330Q (2KQ) were exposed to DOX (1 μM, 6 h). The mRNA transcripts of Bcl-2 were evaluated by qPCR. ** \( P < 0.01 \), 'ns' represents no significance. 18sRNA was used as a reference gene. (E) ChIP analysis was performed in the stably transfected H9C2 cells by using anti-GATA4 antibody in the presence/absence of DOX. GATA4 enrichment on the Bcl-2 promoter region was measured by qPCR. *** \( P < 0.001 \), ** \( P < 0.01 \), 'ns' represents no significance. (F, G) Stably transfected H9C2 cells were exposed to DOX (1 μM) for 24 h and stained with TUNEL or TRITC-phalloidin. Images of the stained cells were captured. The quantification of the number of positively TUNEL stained H9C2 cells (F). Cell area was measured and quantified (G). Scale bar, 20 μm. *** \( P < 0.001 \), 'ns' represents no significance. (H) Cell colony formation showing the survival fraction of H9C2 cells after the indicated doses of DOX for 1 h. The plates (\( n = 3 \)/group) were stained with crystal violet and the colony numbers were counted. ** \( P < 0.01 \), *** \( P < 0.001 \), 'ns' represents no significance.
Figure 6. The SIRT6–TIP60–GATA4 axis facilitates chromatin accessibility. (A, B) ChIP assay was performed using anti-TIP60 or anti-H3K9ac antibody in WT and SIRT6 KO HEK293 cells with or without overexpression of FLAG-TIP60. The enrichment of TIP60 and H3K9ac on Bcl-2 promoter was analyzed by qPCR. **P < 0.01, ***P < 0.001. (C) ChIP analysis was performed using anti-H3K9ac and H3K56ac antibodies in HEK293 cells with or without FLAG-SIRT6 overexpression, showing enrichment of H3K9ac and H3K56ac on the P1 motif of the Bcl-2 promoter region. 'ns' indicates no significance. (D–F) ChIP analysis was performed to test the enrichment of H3K9ac, SIRT6 and TIP60 on the Bcl-2 promoter region by using anti-H3K9ac and antibodies in HEK293 cells with si-NC or si-GATA4. *P < 0.05. 'ns' indicates no significance. (G) Co-IP using anti-FLAG agarose was performed in HEK293 cells expressing HA-GATA4 alone, or with wild-type FLAG-SIRT6 or the FLAG-SIRT6 H133Y mutant. HA-GATA4 levels were analyzed in FLAG-SIRT6 precipitates by western blotting. (H) Deacetylation activity of recombinant SIRT6 after pre-incubation with or without purified His-GATA4 in an in vitro assay. ***P < 0.001. (I) Deacetylation activity of recombinant SIRT6 WT and SIRT6 H133Y after pre-incubation with indicated doses of G4-13 peptides (0, 0.2, 2, 20, 200 μM) in an in vitro assay. ***P < 0.001. 'ns' indicates no significance.
either dimerization of GATA4 is not required for the transcription activity, which is different from GATA3, or the 2KR mutation loses the capacity of dimerization with another 2KR or with WT GATA4. These hypotheses require more evidences to support in future study.

TIP60 belongs to the MYST family of acetyltransferases. Heart-specific depletion of Tip60 increases myocyte density and apoptosis and shortens lifespan in mice (48); however, the underlying mechanism remains to be fully elucidated. Here, we identify TIP60 as a novel acetyltransferase for GATA4 acetylation on K328/330 and protects against stress-induced heart failure. The data provide an explanation by which TIP60 maintains heart homeostasis and a potential target for the prevention of chemotherapy side effects.

In summary, we identify a novel Sirt6–Tip60–GATA4 axis that regulates anti-apoptotic pathway and thus prevent DOX cardiotoxicity. Targeting this axis might constitute a potential target for the prevention of chemotherapy side effects.

SUPPLEMENTARY DATA
Supplementary Data are available at NAR Online.

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