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Permalink
https://escholarship.org/uc/item/5vw7c6mc

Journal
Epigenetics, 7(7)

ISSN
1559-2294

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Publication Date
2012

DOI
10.4161/epi.20733

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Peer reviewed
SIRT1-mediated deacetylation of MeCP2 contributes to BDNF expression

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Keywords: MeCP2, SIRT1, Rett syndrome, acetylation, sirtuins

Methyl-CpG binding protein 2 (MeCP2) binds methylated cytosines at CpG sites on DNA and it is thought to function as a critical epigenetic regulator. Mutations in the MeCP2 gene have been associated with Rett syndrome, a human neurodevelopmental disorder. Here we show that MeCP2 is acetylated by p300 and that SIRT1 mediates its deacetylation. SIRT1, the mammalian homologue of Sir2 in yeast, is a nicotinamide-adenine dinucleotide (NAD+)-dependent histone deacetylase that belongs to the family of HDAC class III sirtuins. Importantly, SIRT1 has been shown to play a critical role in synaptic plasticity and memory formation. This study reveals a functional interplay between two critical epigenetic regulators, MeCP2 and SIRT1, which controls MeCP2 binding activity to the brain-derived neurotrophic factor (BDNF) promoter in a specific region of the brain.

Introduction

Methyl-CpG binding protein 2 (MeCP2) is a very abundant protein that was identified for its capacity to bind methylated cytosines at CpG sites.1-3 MeCP2 is highly conserved within mammals and it contains a N-terminal methyl-CpG-binding domain (MBD) and a C-terminal transcriptional repressor domain (TRD).4 A large body of evidence has established the concept that DNA methylation is associated with transcriptional silencing. The family of MBD binding proteins has been intimately linked to this process.4 Moreover, MeCP2 is capable of further inhibiting transcription through the recruitment of Sin3A and histone deacetylases.5 MeCP2 expression is present in many different tissues, but it appears to be most abundant in the brain.3 The MeCP2 gene, located on the X-chromosome, is transcribed into two different splicing isoforms, MeCP2e1 and MeCP2e2, that differ only in a few amino acids of their N-terminal regions.6 While the e2 isoform was the first one to be identified, the e1 isoform is the most expressed in the brain in both human and mice.7,8 Mutations in the MeCP2 gene have been associated with Rett syndrome,9-10 a neurodevelopmental disorder with affected individuals displaying autistic features, mental retardation and motor and respiratory abnormalities.11 MeCP2 is capable of controlling the expression of several genes that play a crucial role during the process of synapse formation, such as brain-derived neurotrophic factor (BDNF), inhibitors of differentiation (IDs), early growth gene response 2 (EGR2) and JunB.12-14 The molecular and physiological pathways controlling MeCP2 function have remained elusive.

SIRT1, the mammalian homolog of Sir2 in yeast, is a nicotinamide-adenine dinucleotide (NAD+)-dependent histone deacetylase.15 Its function is tightly coupled to cellular metabolism and its dysfunction has been linked to inflammation, obesity and cancer.16 Accumulating evidence underscores the importance of SIRT1-mediated epigenetic control in neuronal plasticity.16,17 It has been described that SIRT1 can regulate neuronal differentiation18,19 and also prevents neurodegeneration in mouse models of Alzheimer disease.20-22 Moreover, Tau acetylation can be reverted by SIRT1.23 Interestingly, SIRT1 brain-specific knock-out mice show impaired cognitive abilities.24 This enzyme can also regulate synaptic plasticity and memory formation through a microRNA-mediated mechanism.25

In this study, we show that MeCP2 is acetylated at a specific lysine residue and that SIRT1 mediates its reversible deacetylation. This post-translational modification appears to modulate MeCP2 binding at the BDNF promoter in the hippocampus. Thus, we reveal the convergence in the control of SIRT1 and MeCP2, two critical epigenetic regulators.

Results

MeCP2 protein is acetylated by p300. Through a high-resolution mass spectrometry screen, it was found that the MeCP2 protein is acetylated at a single site corresponding to lysine 461 of human MeCP2.26 In order to investigate if acetylation could operate as a post-translational modification that regulates MeCP2 function, we first determined if this lysine was conserved between different species. As shown in Figure S1A, the putative acetylated lysine is conserved across several species, and it corresponds to lysine 464 of the mouse e1 isoform, which is the most expressed MeCP2 isoform in the brain. We found that the MeCP2e1 isoform is readily acetylated in vitro (Fig. 1A). To determine whether MeCP2
is acetylated in vivo, we generated a polyclonal antibody that specifically recognizes the acetylated form of MeCP2e1 when modified at lysine 464. We ectopically expressed MeCP2e1 or a mutant with the single amino acid K464R conversion in HEK-293 cells. The K464R mutation mimics a deacetylated lysine residue and is not recognizable by the anti-acetyl MeCP2 antibody (Fig. S1B). Analyzing the primary sequence of the MeCP2e1 protein, we detected several p300 binding motifs.27 We then sought to investigate if this acetyltransferase could mediate MeCP2 acetylation in vitro. We co-transfected p300 or CBP together with MeCP2e1 or the MeCP2e1-K464 point mutant in HEK-293 cells. Co-expression of p300 leads to an efficient increase in the levels of MeCP2 acetylation at lysine 464 (Fig. 1B and C). On the contrary, CBP is not able to efficiently induce acetylation of MeCP2. It should be noted that, even when using equivalent expression vectors, CBP is always expressed at lower levels when compared to p300 (Fig. 1B). To definitely demonstrate that MeCP2 is acetylated, we purified mouse recombinant MeCP2 with glutathione S-transferase (GST) or a control GST-empty, and MeCP2 acetylation was detected only in the presence of purified recombinant p300 (Fig. 1D). In addition, in vitro incorporation assays using 14C-acetyl-Coenzyme A demonstrated that the lysine 464 is acetylated upon addition of p300 (Fig. 1E).

**SIRT1 mediates MeCP2 deacetylation.** The evidence that MeCP2 undergoes acetylation implies that a deacetylase may reverse this post-translational modification. SIRT1 involvement in cognition, memory formation and neurodegenerative disorders has recently been described in references 17, 19, 20 and 24. In addition, SIRT1 has been shown to readily deacetylate and interact with various non-histone substrates.15,17 Thus, we sought to investigate whether SIRT1 interacts with MeCP2. When co-expressed in HEK-293 cells, MeCP2 readily co-immunoprecipitates with SIRT1 (Fig. 1F and G). Strikingly, MeCP2 does not interact with the SIRT1 mutant H363Y, in which a single
This result confirms the capacity of SIRT1 to deacetylate MeCP2 at the targeted lysine in vitro.

MeCP2 acetylation and its effect on BDNF expression. To determine whether MeCP2 acetylation could be detected in vivo, primary cortical neurons were prepared from newborn mice. After 5 d in culture (DIV5), immunofluorescence experiments using the anti-acetyl MeCP2 antibody were performed. As shown in Figure 2B, a clear nuclear signal was detected, demonstrating that acetylation of MeCP2 at K464 occurs in vivo. Importantly, treatment with the SIRT1-specific inhibitor EX527 increased acetylation significantly (~50%) (Fig. 2C). The total amount of MeCP2 protein remains unchanged after the treatment with EX527 (Fig. 2D). These results confirmed that acetylation of MeCP2 is controlled by SIRT1. Consequently, we sought to investigate whether MeCP2 acetylation could modify its recruitment to its primary target, the BDNF promoter. Chromatin immunoprecipitation (ChIP) analyses performed on fresh hippocampus tissue showed that MeCP2 recruitment on the BDNF exon 4 promoter was significantly higher in SIRT1Δ ex4 mice as compared to the wild type littermates (Fig. 3A). This difference in chromatin recruitment was associated to a decrease in both
were purified from wild type pups (1–2 postnatal days). Purified mice were kindly provided by Dr. L.H. Tsai.

**Chromatin immunoprecipitation (ChIP).** Hippocampi were dissected from Nestin-Cre brain and disuccinimidyl glutarate (DSG) was added to a final concentration of 2 mM for crosslinking. After 45 min at room temperature, formaldehyde was added to a final concentration of 1% (v/v) and cells incubated for 15 min. After dual crosslinking, glycine was added to a final concentration of 1% (v/v) and cells quenched formaldehyde. Samples were homogenized, resuspended to a final concentration of 0.1 M and incubated for 10 min to quench formaldehyde. The whole-cell extract was incubated overnight at 4°C with the appropriate antibody. Protein G/salmon sperm was added and after two hours beads were incubated over-night at 4°C with the appropriate antibody. Bound complexes were eluted from the beads with elution buffer (10% SDS, 0.1 mM EDTA, 0.1 M NaCl) in the presence of 0.1 mg recombinant p300 catalytic domain (Active Motif). The reaction was incubated for 1 h at 37°C. Reactions were stopped after 1 h, and the samples were resolved by SDS PAGE and analyzed by western blotting.

**In vitro acetylation assay.** GST-MeCP2e1 was expressed in E. coli BL21. Recombinant proteins were lysed in lysis buffer (20 mM Tris-HCl pH = 8, 0.3 mM EDTA, 20% Glycerol, 5 mM DTT, 0.5 mM PMSF, 1% Triton X-100, 500 mM KCl) and purified by glutathione Sepharose 4B (Amersham). The purified protein was incubated in acetylation buffer (50 mM Tris-HCl pH = 8, 50 mM Acetyl-CoA (SIGMA), 0.1 mM EDTA, 1 mM DTT, 10% glycerol) plus 0.1 mg purified recombinant p300 catalytic domain (Active Motif). The reaction was incubated for 1 h at 37°C. Reactions were stopped by the addition of 2x Laemmli loading buffer, followed by SDS PAGE and western blot analysis. Purified GST-MeCP2e1 was incubated in acetylation buffer (50 mM Tris-HCl pH = 8, 10% glycerol, 100 mM NaCl, 1 mM DTT, 0.2 mM EDTA, 0.2 mm PMSF, 1 mCi of [3H]Acetyl-CoA) in the presence of 0.1 mg recombinant p300 (Active Motif). The reaction was stopped after 1 h, and the samples were resolved by SDS-PAGE. Gels were stained with Coomassie blue, destained, dried and the level of acetyl MeCP2 was detected by autoradiography.

**Antibodies and western blot.** Antibody against FLAG (M2) was from SIGMA, the anti-MeCP2 antibody was from Cell Signaling (MeCP2 D4FS XPTM) and BDNF antibody from Santa Cruz (BDNF N20 sc-546). c-Myc was from Millipore (clone 9E10). Pan-acetyl lysine antibody (Cell Signaling #9441). The polyclonal acetyl-lysine 464 MeCP2e1 was generated by immunizing rabbits with KHL-conjugates of the peptide NH2-AEK(ac) YKHRGEGE (ABE28Millipore). Specificity of the antibody was validated both in vitro and in vivo by performing the appropriate controls, both in our laboratory and by Millipore/Merck. All western blots were visualized using a chemiluminescence detection kit (Perkin-Elmer). At least three independent experiments were performed. Densitometry analysis of the film was performed using Adobe Photoshop.

**Immunofluorescence and quantification.** Cells were fixed using 4% paraformaldehyde in PBS. Blocking was performed using 3% BSA diluted in PBS for 30 minutes at room temperature. Primary antibody used: anti-acetyl MeCP2 (ABE28 Millipore 1:400 dilution) was incubated overnight. Secondary antibody was conjugated with Alexa-488 goat anti-rabbit (Invitrogen). Nuclei were stained with DRAQ-5 [(dimethylamino)-ethylamino-4,8-dihydroxyanthracene-9,10-dione] (Biostatus Limited). Immunolabeled sections were examined with a Leica confocal microscope SP5 (DMRE, Leica). Controls were always performed by omitting primary antibodies. Intensity of fluorescence was evaluated as pixels/μm². The Leica SP5 software LAS AF was used for quantification.

**In vitro deacetylation assay.** Purified MeCP2e1-FLAG was incubated in deacetylation buffer (50 mM Hepes pH = 7.9, 150 mM NaCl, 1 mM DTT) in the presence of purified recombinant human SIRT1 (Sirtris Pharmaceuticals) plus 5 mM NAD+ (SIGMA), 1 mM trichostatin A (TSA), 50 mM EX527 (Tocris), 10 mM nicotinamide (NAM) for 1 h at 37°C. The reactions were resolved on SDS PAGE and analyzed by western blotting.

**Materials and Methods**

**Mice.** The SIRT1Apoe/−Nestin-Cre mice and the SIRT1−/−Nestin-Cre mice were kindly provided by Dr. L.H. Tsai.

**Primary neuronal cultures.** Primary cortical neurons were prepared from wild type pups (1–2 postnatal days). Purified cells were plated using Neurobasal A supplemented with B27 (Invitrogen) on poly-D-lysine coated plates. All the experiments were performed at 5 days in vitro culture (DIV).

**BDNF exon IV-200 FW: 5'-GGC TTC TGT GTG CGT GAA TTT GC-3'; BDNF exon IV 0 REW: 5'-AAA GTG GGT GGG AGT CCA CGA G-3'.**

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**Cell culture, cell extracts and immunoprecipitation.** Human embryonic kidney HEK-293 were maintained at 37°C and 5%
CO₂, in Dulbecco’s modified Eagle’s high glucose (Thermo Scientific) with antibiotics (penicillin and streptomycin) and 10% newborn calf serum (NCS). Cells were washed in phosphate-buffered saline (PBS) and lysed in RIPA buffer [50 mM Tris pH = 8, 150 mM NaCl, 5 mM EDTA, 15 mM MgCl₂, 1% NP40, 1x protease inhibitor cocktail (Roche), 1 mM DTT, 1 mM trichostatin A (TSA), 10 mM NAM, 10 mM NaF, 1 mM PMSF]. Immunoprecipitation was performed by pre-clearing 500 mg-1 mg of whole lysates with protein G-Agarose beads for two hours, and then by incubating with the appropriate amount of antibody or with anti FLAG-M2 Affinity gel at 4°C (SIGMA).

**Plasmids.** MeCP2e1-FLAG pCMV-TAG4 (STRATAGENE) was a kind gift of Dr. J.M. LaSalle. Mouse MeCP2e1 cDNA was cloned in a 6-myc/pCDNA3 plasmid. The point mutant MeCP2e1 K464R was generated using the Agilent’s QuickChange Site-Directed Mutagenesis Kit. The truncation mutants of MeCP2 were generated by PCR amplification followed by cloning in 6-myc/pCDNA3 or pCMV-TAG4 (STRATAGENE). StratageneQuickChange Site-Directed Mutagenesis Kit was used to generate the single point mutant K464R MeCP2e1.

**Statistical analysis.** Differences between two means were assessed with Student’s t-test. At least three independent experiments were performed (n = 3). *p < 0.05 was considered significant, **p < 0.01 and ***p < 0.001 were considered highly significant.

**Conclusion.** Methyl-CpG binding protein 2 (MeCP2) is capable of binding methylated cytosines at CpG sites, and mutations in its gene have been associated with Rett Syndrome. It has been previously described that post-translational modifications can regulate MeCP2 protein function. Phosphorylation at serine 80 or serine 421 regulates, in opposite ways, the binding or release of MeCP2 from BDNF gene promoter. 29,30 In this study, we demonstrated that MeCP2 undergoes lysine acetylation, an event mediated by the acetyltransferase p300 in vitro. More interestingly, SIRT1, a NAD⁺-dependent histone deacetylase, appears to mediate the opposite reversible reaction. Importantly, accumulating evidence underscores the importance of this histone deacetylase in regulating neuronal differentiation. 31,32

We speculate that SIRT1-dependent deacetylation of MeCP2 could allow its release from the methylated CpG sites within the BDNF exon 4 promoter leading to increased BDNF transcription. In keeping with this hypothesis, the absence of a functional SIRT1 in SIRT1Δ4 mice prevents the release of the acetylated MeCP2 from the BDNF promoter, resulting in decreased expression.

**Disclosure of Potential Conflicts of Interest**
No potential conflicts of interest were disclosed.

**Acknowledgments**
We are grateful to Li-Huei Tsai and Janine LaSalle for discussions, help and critical reading of the manuscript. We thank all the members of the Sassone-Corsi laboratory for helpful discussions. This work was supported by the National Institute of Health and Sirtris-GSK. L.Z. was in part supported by the American-Italian Cancer Foundation, New York.

**Supplemental Material**
Supplemental materials may be found here: www.landesbioscience.com/journals/epigenetics/article/20733

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