

KDM3B Is the H3K9 Demethylase Involved in Transcriptional Activation of *lmo2* in Leukemia

Ji-Young Kim,^a Kee-Beom Kim,^a Gwang Hyeon Eom,^b Nakwon Choe,^b Hae Jin Kee,^b Hye-Ju Son,^a Si-Taek Oh,^a Dong-Wook Kim,^c Jhang Ho Pak,^d Hee Jo Baek,^e Hoon Kook,^e Yoonsoo Hahn,^a Hyun Kook,^b Debabrata Chakravarti,^f and Sang-Beom Seo^a

Department of Life Science, College of Natural Sciences, Chung-Ang University, Seoul, Republic of Korea^a; Medical Research Center for Gene Regulation and Department of Pharmacology, Chonnam National University, Gwangju, Republic of Korea^b; Department of Oral Pharmacology, School of Dentistry and Institute of Oral Bioscience, Chonbuk National University, Jeonju, Republic of Korea^c; University of Ulsan College of Medicine, Asan Medical Center, Seoul, Republic of Korea^d; Department of Pediatrics, Chonnam National University Hwasun Hospital, Hwasun, Republic of Korea^e; and Division of Reproductive Biology Research, Department of Obstetrics and Gynecology and Robert H. Lurie Comprehensive Cancer Center, Northwestern University Feinberg School of Medicine, Chicago, Illinois, USA^f

Histone lysine methylation and demethylation are considered critical steps in transcriptional regulation. In this report, we performed chromatin immunoprecipitation with microarray technology (ChIP-chip) analysis to examine the genome-wide occupancy of H3K9-me2 during all-*trans*-retinoic acid (ATRA)-induced differentiation of HL-60 promyelocytic leukemia cells. Using this approach, we found that KDM3B, which contains a JmjC domain, was downregulated during differentiation through the recruitment of a corepressor complex. Furthermore, KDM3B displayed histone H3K9-me1/2 demethylase activity and induced leukemogenic oncogene *lmo2* expression via a synergistic interaction with CBP. Here, we found that KDM3B repressed leukemia cell differentiation and was upregulated in blood cells from acute lymphoblastic leukemia (ALL)-type leukemia patients. The combined results of this study provide evidence that the H3K9-me1/2 demethylase KDM3B might play a role in leukemogenesis via activation of *lmo2* through interdependent actions with the histone acetyltransferase (HAT) complex containing CBP.

Chromatin remodeling facilitated by various histone modifications is considered an important factor in the transcriptional regulation of target genes. Since the initial identification of Suv39h1, which is a histone lysine methyltransferase (HMTase), numerous other HMTases with different specificities toward histone lysine residues and target genes have been discovered (16, 22, 27). Histone lysine methylation is linked to both activation and repression of chromatin, and its activity is dependent on which lysine is targeted. Among these, methylation of histone H3K9 is one of the most intensively studied histone modifications. Histone H3K9 methylation by Suv39h1 is followed by binding of heterochromatin protein 1 (HP1) and association with corepressor complexes, defining the fundamental role of H3K9 methylation in heterochromatin formation and silencing of target genes (8, 17). Methylation of H3K9 by Suv39h1, ESET/SETDB1, and G9a further indicates the importance of this event in the silencing and repression of various target genes during disease development and progression (1, 6, 25). Although recent studies have demonstrated the potential role of histone modifiers in the development of leukemia, the underlying molecular mechanisms, such as target gene modulation and diverse interactions by these modifiers, remain to be determined.

Differentiation of the human promyelocytic leukemia cell line HL-60 can be induced toward granulocytes by all-*trans*-retinoic acid (ATRA) or dimethyl sulfoxide (DMSO), toward neutrophils by dimethylformamide (DMF), or toward monocytes/macrophages by phorbol esters, via either a monocytic or granulocytic pathway (3). Acute promyelocytic leukemia (APL) can be effectively treated with differentiation therapy using ATRA, which induces terminal differentiation followed by apoptosis. Although both biochemical and molecular studies have linked ATRA-mediated differentiation in leukemia cell lines with transcriptional regulation, the detailed mechanism

and role of histone modification have not yet been fully determined. Recent studies have suggested that PML/RAR α and PLZF/RAR α fusion proteins recruited polycomb repressive complex 2 (PRC2) to the RAR β 2 promoter (27). Mixed-lineage leukemia (MLL) or the MLL fusion protein has been shown to mediate both positive and negative regulation of target genes through the recruitment of different epigenetic regulators. Interestingly, components of the MLL complex and the H3K27 demethylase UTX complex are recruited to ATRA response genes after ATRA treatment and activate the transcription of Hox genes (19).

To gain further insight into the chromosome-wide dimethylation of histone H3K9 target sites in differentiated leukemia cells after ATRA treatment, we performed a chromatin immunoprecipitation (ChIP) assay using antibodies that recognize dimethylated H3K9. The precipitated DNA was hybridized to a promoter array, which allowed us to analyze the 3,500-bp upstream region and 750-bp downstream region of the transcriptional start site (TSS) of 13,000 human genes. By using this approach, we identified more than 220 target gene promoters, including that of *KDM3B*, which were repressed during ATRA-mediated differentiation.

In this study, we further focused on the JmjC domain-containing histone KDM3B and characterized its transcriptional

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Address correspondence to Sang-Beom Seo, sangbs@cnu.ac.kr.

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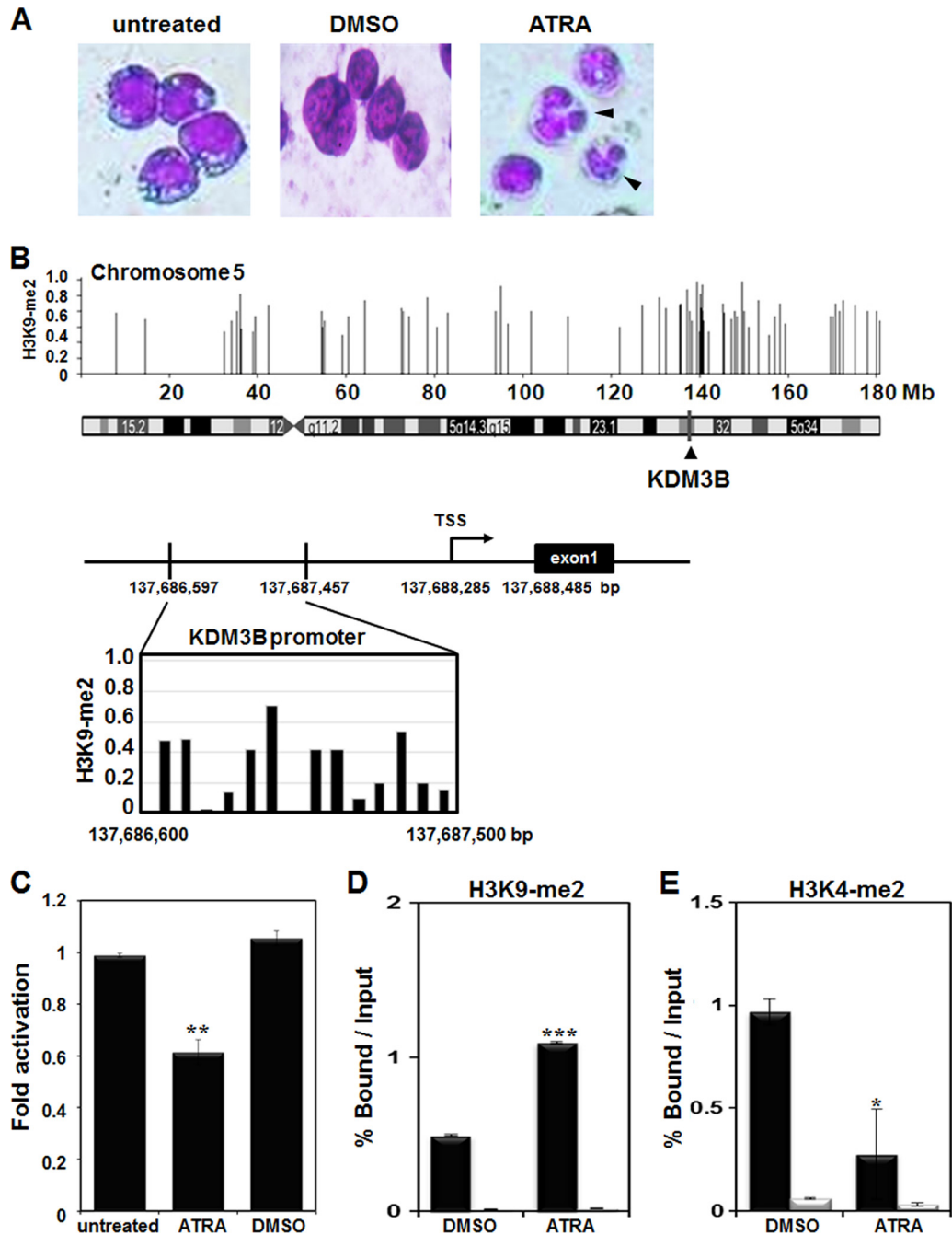


FIG 1 Identification of KDM3B as an H3K9-me2 target gene in ATRA-treated HL-60 cells. (A) Morphological observation of Giemsa-stained cells treated for 48 h with 1 μ M ATRA. (B) The *KDM3B* locus from chromosome 5 is shown in 5' to 3' orientation. The magnified locus of *KDM3B* was highly enriched in H3K9-me2 in ATRA-treated HL-60 cells. (C) The expression level of KDM3B in ATRA-treated HL-60 cells was detected via real-time PCR. (D to G) ATRA-treated HL-60 cells were differentiated for 48 h. Cross-linked samples were immunoprecipitated with anti-H3K9-me2 (D), anti-H3K4-me2 (E), anti-AcH3 and anti-AcH4 (F), or anti-HDAC1, anti-HP1 α , and anti-SMRT (G) antibodies. The precipitated DNA fragments were subjected to real-time PCR in the *KDM3B* proximal or distal promoter region. Normalization was performed using input DNA levels as controls in the same reaction. Results are shown as means \pm SDs; $n = 3$. *, $P < 0.05$; **, $P < 0.01$; ***, $P < 0.001$. (H) A mouse adult tissue blot was hybridized to anti-KDM3B antibodies. GAPDH (glyceraldehyde-3-phosphate dehydrogenase) was used as a control.

regulatory role in leukemia via H3K9-me1/2 demethylase activity. Genome-wide analysis identified a subset of KDM3B target genes, including leukemic oncogene *lmo2*. KDM3B was recruited to the leukemogenic oncogene *lmo2* promoter along with CBP and was found to function as a coactivator of the

target gene. In addition, KDM3B was shown to function as an epigenetic regulator of *lmo2*-mediated leukemogenesis. Furthermore, KDM3B expression induced leukemic transformation by inhibiting leukemia cell differentiation and was upregulated in blood cells obtained from certain types of leukemia patients.

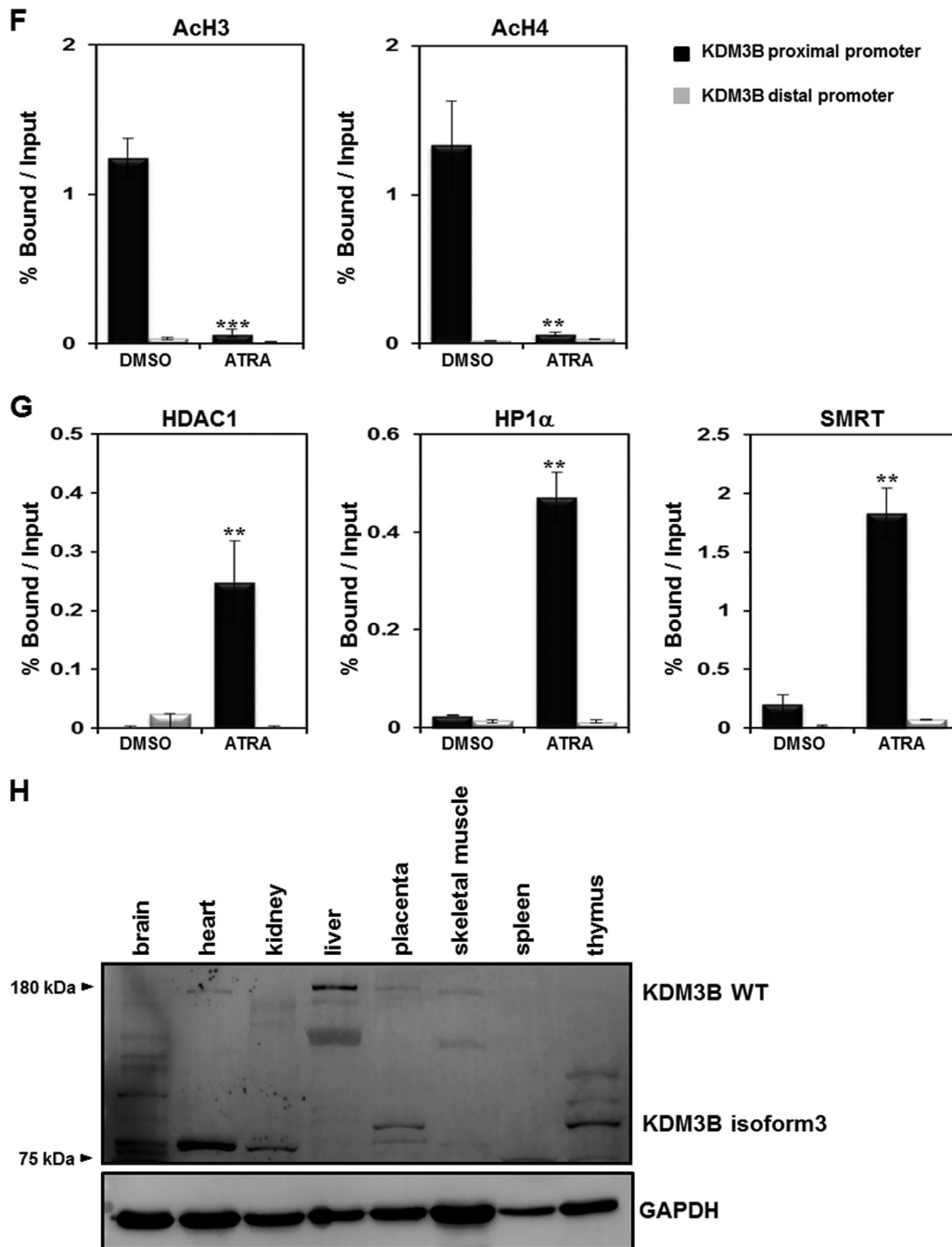


FIG 1 continued

MATERIALS AND METHODS

Plasmid. pOBT7-KDM3B (BC001202, amino acids 1003 to 1761, KDM3B isoform 3) was purchased from Open Biosystems. KDM3B isoform 3 was inserted into the pGEX-4T1 bacterial expression vector to construct GST-KDM3B fusion proteins and mammalian expression vectors. KDM3B (H558A) point mutants were constructed by Cosmo Genetech. In order to construct the mammalian expression vectors, we used modified pcDNA6-HA-myc-his (Invitrogen) and GFP-C1 (Clontech) to create the hemagglutinin (HA)-, myc-, and His-tagged KDM3B and green fluorescent protein (GFP)-tagged KDM3B proteins, respectively. Short hairpin RNA (shRNA) against human *KDM3B* (SHCLNG-

NM-016604) and small interfering RNA (siRNA) against human *KDM3B* (L-020378-01-0010) were purchased from Sigma and Dharmacon, respectively. The effect of KDM3B on P1 promoter-directed expression was evaluated by RNA interference (RNAi) using the Knockout single-vector-inducible RNAi system (Clontech). siRNAs targeting *KDM3B* mRNA were designed by using the siRNA sequence designer software (Clontech). We produced a double-stranded oligonucleotide for shRNA plasmid construction containing the following primers from the 5' to the 3' end: sense primer, 5'-TCGAGGCCTGCAACTTGACTGATACTTCAAGAGAGTATCAGTCAAGTTGCAGGTTTTTACGCGT-3'; antisense primer, 5'-C GGACGTTGAACTGACTATGAAGTTCTCTCATAGTCAGTTCAACG

TCCAAAAAATGCGCATTCGA-3'. This oligonucleotide was cloned into the XhoI-HindIII site of the pSingle-tTS-shRNA vector (Clontech), in which shRNA is expressed under the control of the U6 promoter. siRNA against human *lmo2* (sc-38027) was purchased from Santa Cruz Biotechnology.

Antibodies. Antibodies against KDM3B (X-25, sc-101987; Santa Cruz Biotechnology), histone H3 (06-755; Millipore), *lmo2* (N-16, sc-10497; Santa Cruz Biotechnology), and β -actin (sc-47778; Santa Cruz Biotechnology) were used for immunoblot analysis and the ChIP assay. anti-GFP (B-2, sc-9996; Santa Cruz Biotechnology), CBP (A-22, sc-369; Santa Cruz Biotechnology), and the p300/CBP-associated factor (PCAF) antibody (H-369, sc-8999; Santa Cruz Biotechnology) were used for immunoprecipitation. Antibodies against histone H3K4-me2 (07-030; Millipore), H3K9-me1 (07-450; Millipore)/H3K9-me2 (07-441; Millipore)/H3K9-me3 (17-625; Millipore), H3K27-me2 (07-452; Millipore), and H3K36-me2 (07-274; Millipore) were used for immunoblot analysis or immunocytochemistry.

Stable cell lines. K562 (5×10^6) cells were seeded into a 60-mm dish and were transfected with pSingle-tTS-shRNA-KDM3B using Lipofectamine 2000 (Invitrogen), after which the stably transfected clones were selected in medium containing 2.5 mg/ml of G418 (Sigma). Cells were cultured in RPMI 1640 in the presence or absence of doxycycline (Sigma) and supplemented with 10% of tetracycline (Tet) system-approved fetal bovine serum (FBS) (Clontech), which is a tetracycline-free serum that was developed for the tetracycline-controllable expression system. Forty-eight hours later, RNA extraction and reverse transcription were performed.

Chromatin immunoprecipitation with microarray technology (ChIP-chip) analysis. ChIP was carried out as essentially described in the protocols from Millipore. Briefly, HL-60 cells were treated with 1 μ M ATRA or DMSO and harvested 48 h later. Cells were cross-linked with 1% formaldehyde in the medium at 37°C for 10 min, followed by the addition of 125 mM glycine at room temperature for 5 min, and then scraped into SDS lysis buffer. Samples were further sonicated and diluted for immunoprecipitation with antibodies against H3K9-me2. The immunoprecipitates were eluted and cross-link reversed. DNA fragments were purified using a PCR purification kit (Qiagen). Chromatin-immunoprecipitated DNA was amplified in 2 steps, including a T7 Sequenase extension step using a random primer with a fixed sequence linker and a second step of amplification using the fixed sequence primer and *Taq* polymerase. The products were purified and labeled with amino-allyl-conjugated dUTP using the BioPrime labeling kit (Invitrogen) and a random primer. Cross-linking reactions were quenched with 125 mM glycine, and the Cy3-conjugated DNA and Cy5-conjugated background DNA were mixed and purified with the PCR purification kit. The purified probe was combined with hybridization buffer (5 \times SSC [1 \times SSC is 0.15 M NaCl plus 0.015 M sodium citrate], 25% formamide, 0.1% SDS). The hybridization mix was boiled for 5 min and centrifuged for 2 min before being applied to a prehybridized microarray. Microarrays were prehybridized with 5 \times SSC, 25% formamide, 0.1% SDS, and 1% bovine serum albumin (BSA) for 45 min at 42°C. Slides were hybridized for 12 to 16 h at 42°C and washed according to the manufacturer's conditions. ChIP-chip analysis was performed using oligonucleotide-based promoter tiling arrays from NimbleGen. Briefly, microarrays were scanned with an Axon 4000A scanner (Axon Instruments), and images were analyzed with SignalMap software. The data were filtered to remove spots with aberrant morphology or those with intensities below the threshold of detection, and then median ratios were normalized to background.

Immunocytochemistry. 293T cells were grown on chamber slides and transfected with GFP-KDM3B expression plasmids using Lipofectamine 2000 (Invitrogen). After 48 h of transfection, the cells were fixed with 4% paraformaldehyde and incubated with each antibody, washed, and stained with SYTO61 (Invitrogen). The slides were then mounted with Gel/Mount (BioMeda), and images were acquired with a confocal laser scanning microscope (Zeiss LSM5 Exiter).

Demethylase assay and mass spectrometry. Demethylase assays were conducted as described previously (31). H3K9-me1 peptide [TARK(me1)STG], H3K9-me2 peptide [TARK(me2)STG], and H3K9-me3 peptide [TARK(me3)STG] were synthesized based on the N-terminal amino acid sequences of histone H3 (Peptron). Synthetic peptides (0.1 μ g) were used as the substrates in the demethylase assay with GST-KDM3B. The reaction was stopped by 10% trichloroacetic acid (TCA) precipitation. After removing the precipitates by centrifugation, the supernatants were retrieved, and methylated peptides in the supernatants were analyzed by liquid chromatography-mass spectrometry (LC-MS) at the Korea Basic Science Institute.

Transcriptional activity assay. For the transcriptional activity assay, 293T cells were seeded at 3×10^4 cells/well in a 48-well plate and were cotransfected with 150 to 300 ng of the expression plasmid and 150 ng of the pGL2-*lmo2* promoter (−661 to −330) reporter plasmid using Lipofectamine 2000. After 48 h, the cells were harvested and subjected to a luciferase assay (Promega). The level of β -galactosidase activity was used to normalize reporter luciferase. Data are expressed as the means of four replicates from a single assay. The results shown are representative of at least three independent experiments.

Immunoprecipitation. For the interaction assays, 293T cells were cotransfected with pcDNA6-KDM3B, Gal4-CBP, or FLAG-PCAF, lysed in radioimmunoprecipitation assay (RIPA) lysis buffer, and then immunoprecipitated with anti-KDM3B, anti-CBP, or anti-PCAF antibodies and protein A/G-agarose beads (GenDEPOT). The bound proteins were analyzed in immunoblots with anti-KDM3B, anti-CBP, and anti-PCAF antibodies.

Reverse transcription and real-time PCR. Total RNA samples from the KDM3B- or sh-KDM3B construct-transfected HL-60 and 293T cells were extracted using RNAsiso Plus (TaKaRa) in accordance with the manufacturer's recommendations. Total RNA (2 μ g) was used to synthesize cDNA. cDNA synthesis was primed using an oligo(dT) primer (Fermentas), and the quantified cDNA was used for *KDM3B*, *lmo2*, *Raf-1*, and *Myc* mRNA expression pattern analysis. The primer sequences were as follows: for *lmo2*, 5'-GCGCGGTGACTGTCCTTGGAG-3' (sense) and 5'-GTCTCTCCGGCAGAGCTTCC-3' (antisense); for *Raf-1*, 5'-AGAGGTGATCCGAATGCAGGA-3' (sense) and 5'-AGCAGCTCAATGGAAGACAGG-3' (antisense); and for *Myc*, 5'-TACCCTCTCAACGACAGCAG-3' (sense) and 5'-TCTTGACATTCTCCTCGGTG-3' (antisense). The amplification reaction was performed under the following conditions: 40 cycles of denaturation at 94°C for 15 s, annealing at 60°C for 30 s, and extension at 72°C for 30 s. Dissociation curves were generated after each PCR run to ensure that a single product of the appropriate length was amplified. The mean threshold cycle (C_T) and standard error were calculated from individual C_T values obtained from three replicates per stage. The normalized mean C_T was estimated as ΔC_T by subtracting the mean C_T of β -actin from those of *KDM3B* and *lmo2*. $\Delta\Delta C_T$ was calculated as the difference between the control ΔC_T and the values obtained for each sample. The fold change in gene expression, relative to the untreated control, was calculated as $2^{-\Delta\Delta C_T}$.

Microarray analysis. For KDM3B target gene profiling, we used the Illumina HumanHT-12-v3-BeadChip (Illumina), which includes a bead pool of more than 48,000 unique bead types corresponding to 48,804 transcripts. Total RNA (500 ng) isolated from K562 cells stably expressing short hairpin RNA control (sh-CTL) and sh-KDM3B were reverse transcribed and amplified according to the protocols described in the Illumina TotalPrep RNA amplification kit manual (Ambion). *In vitro* transcription was then carried out to generate cRNA. cRNA (750 ng) was hybridized onto each array (two replicates for each condition) and then labeled with Cy3-streptavidin (Amersham Biosciences). The array was then scanned using a BeadStation 500 system (Illumina). The full data set was submitted to Gene Expression Omnibus under submission number GSE30294. The probe intensities were normalized using quantile normalization in BeadArray 1.10.0, an R/Bioconductor package. The probes were annotated using lumi 1.8.3, an R/Bioconductor package.

TABLE 1 Identification of H3K9-me2 target genes during differentiation of a leukemia cell line by ATRA

Gene name	GenBank accession no.	Description	H3K9-me2 enrichment	FDR ^a	TSS (bp)	Distance (bp)
<i>OR51E1</i>	BC022401	Olfactory receptor, family 51, subfamily E, member 1 gene	0.72	0	4,621,731	-2,060
<i>ALCAM</i>	AK127617	Activated leukocyte cell adhesion molecule gene	0.71	0	106,735,135	-2,170
<i>NRG2</i>	NM_004883	Neuregulin 2 gene	0.59	0	-139,403,063	-1,877
<i>ANK3</i>	BX648574	Ankyrin 3 gene, node of Ranvier (ankyrin G)	0.59	0.02	-62,163,254	-2,974
<i>NUP210L</i>	NM_207308	Nucleoporin 210 kDa-like gene	0.57	0.019	-152,378,952	-1,681
<i>EBF2</i>	AY700779	Early B cell factor 2 gene	0.57	0.019	-25,801,349	472
<i>CREBBP</i>	U85962	CREB-binding protein gene	0.57	0.168	-3,870,723	-1,613
<i>JAK2</i>	NM_004972	Janus kinase 2 (a protein tyrosine kinase) gene	0.47	0.092	4,975,244	-1,772
<i>JAKMIP2</i>	NM_014790	Janus kinase and microtubule interacting protein 2 gene	0.35	0.013	-147,142,445	-1,102
<i>KDM3B</i>	NM_016604	Lysine (K)-specific demethylase 3B gene	0.4	0.024	137,688,285	-1,258

^a FDR, false discovery rate.

ChIP assay and real-time PCR. HL-60 and 293T cells were transfected with KDM3B or sh-KDM3Bs, harvested after 48 h, and then cross-linked with 1% formaldehyde added to the medium for 10 min at room temperature, followed by the addition of 125 mM glycine for 5 min at room temperature. The samples were sonicated, and the lysates were subjected to immunoprecipitation using the indicated antibodies. The immunoprecipitates were eluted and reverse cross-linked, after which the DNA fragments were purified for PCR amplification. To analyze the *KDM3B* promoter region, primer sets consisting of the 457-bp region (nucleotides -1067 to -611; sense, 5'-CACAGAAGCCTTGGAGAAAGAACC-3', and antisense, 5'-AAGTCTACGTGTTCTCCATTC-3') or the *KDM3B* distal promoter region (nucleotides -3036 to -2884; sense, 5'-CCCCACAAAGACAAGGAAGA-3', and antisense, 5'-AAGCCTGGGC CACAGTATAA-3') were used. A 488-bp fragment corresponding to nucleotides -687 to -200 of the *lmo2* promoter and the 446-bp fragment (+243 to +688) of the *lmo2* exonic region were PCR amplified. The primer sequences were as follows: for the *lmo2* promoter, 5'-GGACCTG GGACCTCGAAC-3' (sense) and 5'-GGGCTTCTCTCTCGGGAA G-3' (antisense), and for the *lmo2* exonic region, 5'-GCGCGGTGACTG TCCTTGAG-3' (sense) and 5'-GTCTCTCCGGCAGAGCTCC-3' (antisense). For the analysis of the *Myc* and *Raf-1* promoter regions, the primer sets consisted of the 248-bp fragment (nucleotides -710 to -463; sense, 5'-TAATCATTCTAGGCATCGTTTT-3', and antisense, 5'-ATCA TCGCAGCGGAACAGCTG-3') and the 180-bp fragment (nucleotides -430 to -251; sense, 5'-TGCCTCAGCCTCCCGAGTAGCTGGGAC-3', and antisense, 5'-ACTTCCATTAATATTTTCATAGATGAGG-3'), respectively. The primer concentration for the real-time PCR was 0.2 μM/25 μl. The thermal cycler conditions were as follows: 15 min of holding at 95°C, followed by 45 cycles at 94°C for 15 s, 60°C for 30 s, and 72°C for 30 s (Bio-Rad).

Evaluation of myeloid differentiation. HL-60 cells transfected with KDM3B or sh-KDM3B were treated with 1 μM ATRA. Granulocyte differentiation was evaluated 72 h after ATRA induction based on cell morphology on Wright-Giemsa-stained slides. The expression of CD11b and CD14 was detected 72 h after induction by fluorescence-activated cell sorter (FACS) analysis. For the FACS assay, the cells were stained with the CD14-allophycocyanin (APC) (eBioscience) and CD11b-phycoerythrin (PE) (Millipore) antibodies, which were measured using FACSAria I (BD Biosciences) at the National Instrumentation Center for Environmental Management (NICEM). The results were analyzed using the FACScan Cell Quest software (BD Biosciences).

Leukemia patient samples. Mononuclear cells were obtained from the bone marrow of 30 leukemic patients at the time of diagnosis. Fourteen patients had precursor B cell-type acute lymphoblastic leukemia (ALL), six patients had TEL/AML1 ALL, three patients had early precursor B cell-type ALL, three patients had T cell-type ALL, four patients had AML-ETO-1-type ALL, and nine patients had acute myeloid leukemia

(AML). Informed consent was obtained from the guardians of each patient, and study approval was obtained from the Institutional Review Board of Chonnam National University Hospital.

Oncomine database analysis. The Oncomine database was used as previously described (23). Gene expression data of KDM3B were retrieved from the Oncomine website (www.oncomine.org). Data sets with differences in KDM3B expression between normal and cancer tissues were selected.

Statistical analysis. Data are expressed as the means and standard deviations (SDs) of three or more independent experiments. Statistically significant effects ($P < 0.05$) were evaluated with Microsoft Excel. Differences between groups were evaluated by one-way analysis of variance (ANOVA), followed by Student's *t* test or Bonferroni's test as appropriate.

RESULTS

Identification of the H3K9-me2 target gene *KDM3B* during differentiation of a leukemia cell line by ATRA. To identify direct targets of H3K9-me2 during differentiation in leukemia cells, we performed ChIP-chip analysis using a human promyelocytic leukemia cell line, HL-60, after treatment with ATRA. Differentiation of HL-60 cells by ATRA treatment was confirmed by Giemsa staining (Fig. 1A). Target TSS regions identified by ChIP-chip analysis are listed in Table 1. Among the profiled target TSS regions, we focused on *KDM3B* as a potential target gene of H3K9-me2 modification during leukemia cell differentiation. *KDM3B* is located on human chromosome 5 (5q31), and the target site of H3K9-me2 was located in the *KDM3B* promoter, indicating that modification may have occurred during differentiation (Fig. 1B). Full-length human KDM3B is 1,761 amino acids long and has 2 isoforms by alternative splicing. Isoform 2 is 1,417 amino acids long with the N-terminal amino acids 1 to 344 missing, and isoform 3 is 759 amino acids long with the N-terminal amino acids 1 to 1002 missing.

Verification of the H3K9 target gene *KDM3B* promoter. HL-60 cells were treated with ATRA, and expression of KDM3B and recruitment of chromatin remodelers to the *KDM3B* promoter were analyzed. First, we verified that KDM3B expression was downregulated and that the *KDM3B* promoter was enriched with H3K9-me2 during ATRA-mediated differentiation of HL-60 cells (Fig. 1C and D). On the other hand, H3K4-me2 levels in the *KDM3B* promoter decreased during differentiation (Fig. 1E). As expected, histone acetylation of H3 and H4 significantly decreased when cells were treated with ATRA (Fig. 1F). Since histone modifications, such as H3K9-me2, can recruit various chromatin re-

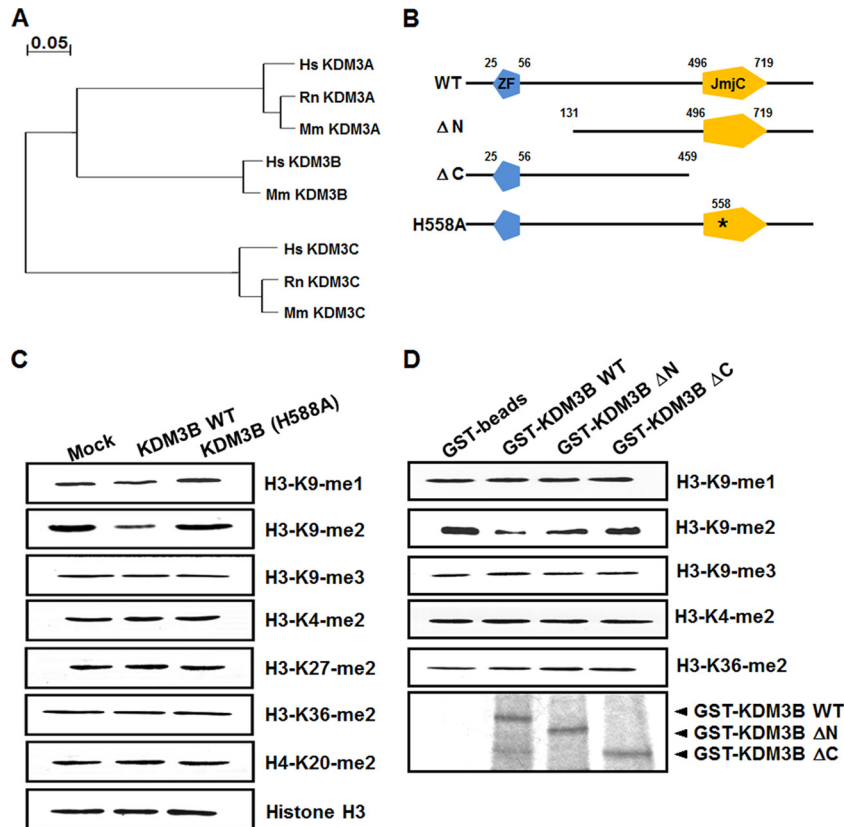


FIG 2 KDM3B is an H3K9-specific demethylase. (A) Phylogenetic analysis based on protein sequence data of human, rat, and mouse KDM3 families. (B) Schematic representations of KDM3B with functional domains and the deletion mutants are shown. (C) 293T cells were transfected with KDM3B and KDM3B (H558A) point mutants, and histone demethylase activities were analyzed using immunoblot analysis with the indicated antibodies. (D) *In vitro* demethylase assays with GST-KDM3B or GST-KDM3B deletion mutants were conducted with core histones and then immunoblotted against the indicated antibodies. (E and F) 293T cells were transfected with GFP-KDM3B or GFP-KDM3B (H558A) and immunostained with antibodies against the indicated histone H3 lysine residues with different methylation marks. SYTO61 staining represents nucleic acids. Arrowheads indicate transfected cells. (G) H3K9-me1/2/3 peptides were incubated with GST-KDM3B, GST-KDM3B (H558A), or GST, and the modified peptides were analyzed via LC-MS.

modeling corepressors that alter the transcription of target genes, we examined the association between cofactors in the *KDM3B* promoter. As expected, enhanced association between the corepressors HDAC1, HP1 α , and SMRT and the *KDM3B* promoter after ATRA treatment was observed by the ChIP assay and real-time PCR (Fig. 1G). Increased binding of this corepressor complex to the *KDM3B* promoter further confirmed that KDM3B was highly repressed during HL-60 cell differentiation induced by ATRA treatment. Using tissue blot analysis with multiple mouse tissue samples, we found that wild-type KDM3B was abundant in the liver, heart, and placenta whereas KDM3B isoform 3 was highly expressed in the placenta and thymus, a finding which implies the possible role of KDM3B in hematopoiesis (Fig. 1H).

KDM3B is a novel histone H3K9 demethylase. KDM3B belongs to a subfamily of proteins containing a JmjC domain that includes KDM3A (JHDM2A) and KDM3C (JHDM2C) (Fig. 2A). These proteins possess a protein domain structure containing a zinc finger (ZF)-like domain at their amino terminus and a JmjC domain near their carboxy terminus. A recent study showed that KDM3C demethylates H3K9-me1/2, indicating that KDM3B is a candidate H3K9 demethylase (15). To verify that KDM3B also possesses H3K9 lysine specificity, KDM3B was overexpressed in 293T cells, and histone modification was

analyzed using specific antibodies. The results of this analysis showed that KDM3B specifically demethylated H3K9-me2 (Fig. 2C). No KDM3B demethylase activity on H3K9-me1 was observed in cells transfected with a KDM3B point mutant (H558A), in which a conserved histidine in the JmjC domain was replaced by an alanine (Fig. 2B and C). Similar results were obtained in an *in vitro* demethylase assay using glutathione *S*-transferase (GST)-purified KDM3B and core histones. Deletion of either the ZF-like motif (KDM3B Δ N) or the JmjC domain (KDM3B Δ C) resulted in a loss of H3K9-me2 demethylase activity (Fig. 2B and D). Next, we used immunofluorescence to analyze H3K9 demethylation by KDM3B. As was reported in other studies that examined KDM3B substrate specificity, we observed a significant loss of H3K9-me1/2 methylation in cells expressing KDM3B (Fig. 2E). However, an amino acid substitution in the JmjC domain (KDM3B H558A) of KDM3B completely abrogated its demethylase activity (Fig. 2F).

Synthetic H3K9-me1/2/3 peptides were incubated alone or in the presence of KDM3B and analyzed by mass spectrometry. In this analysis, the H3K9-me1/2 peptides were shown to be converted into mono- and unmethylated peptides after incubation with KDM3B (Fig. 2G). No demethylase activity was detected when H3K9-me3 peptides were incubated with KDM3B (Fig. 2G).

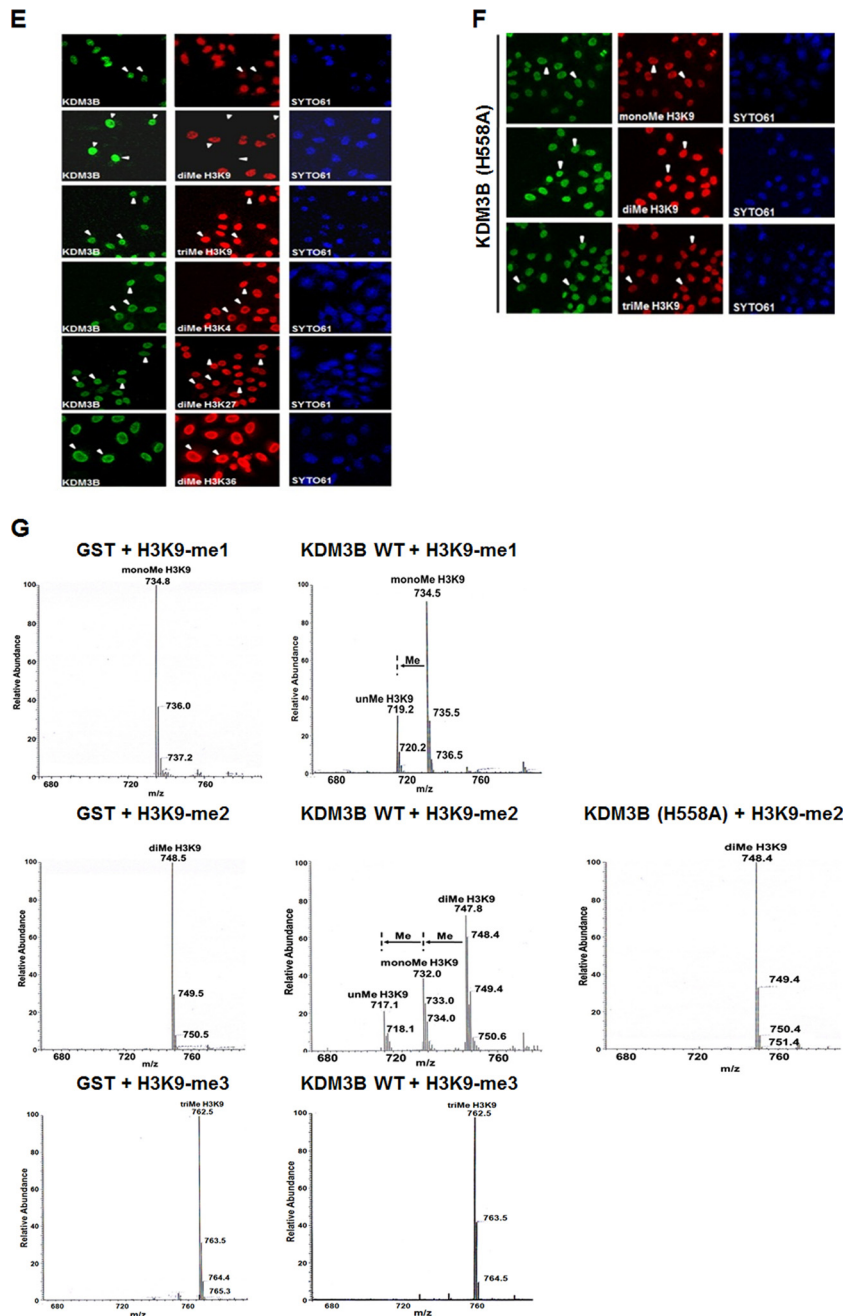


FIG 2 continued

Western blot analysis showed that KDM3B had weak demethylation activity toward H3K9-me1, whereas mass spectrometry and immunofluorescence demonstrated that KDM3B could remove monomethyl groups from the peptides. This most likely occurred because a reduced level of monomethylation was masked by the production of monomethylated histones from a dimethylated histone by KDM3B.

KDM3B regulates the transcription of the *lmo2* oncogene. Since methylation of histone H3K9 has been reported to be involved in transcriptional repression, the demethylation of H3K9 by KDM3B indicates that KDM3B may play a role in transcrip-

tional activation. Having established that KDM3B possesses both H3K9-me1/2 demethylase and transcriptional activation activities, we next searched for genes directly targeted by KDM3B. Downregulation of KDM3B gene expression during ATRA-induced differentiation of leukemia cells was in contrast to its abundant expression in tissues involved in hematopoiesis. This prompted us to focus on several leukemia-related oncogenes as potential candidate target genes of KDM3B. Specifically, the leukemic oncogene *lmo2* showed decreased expression during ATRA-induced differentiation (Fig. 3A). On the other hand, the expression of other genes that play roles in similar oncogenic

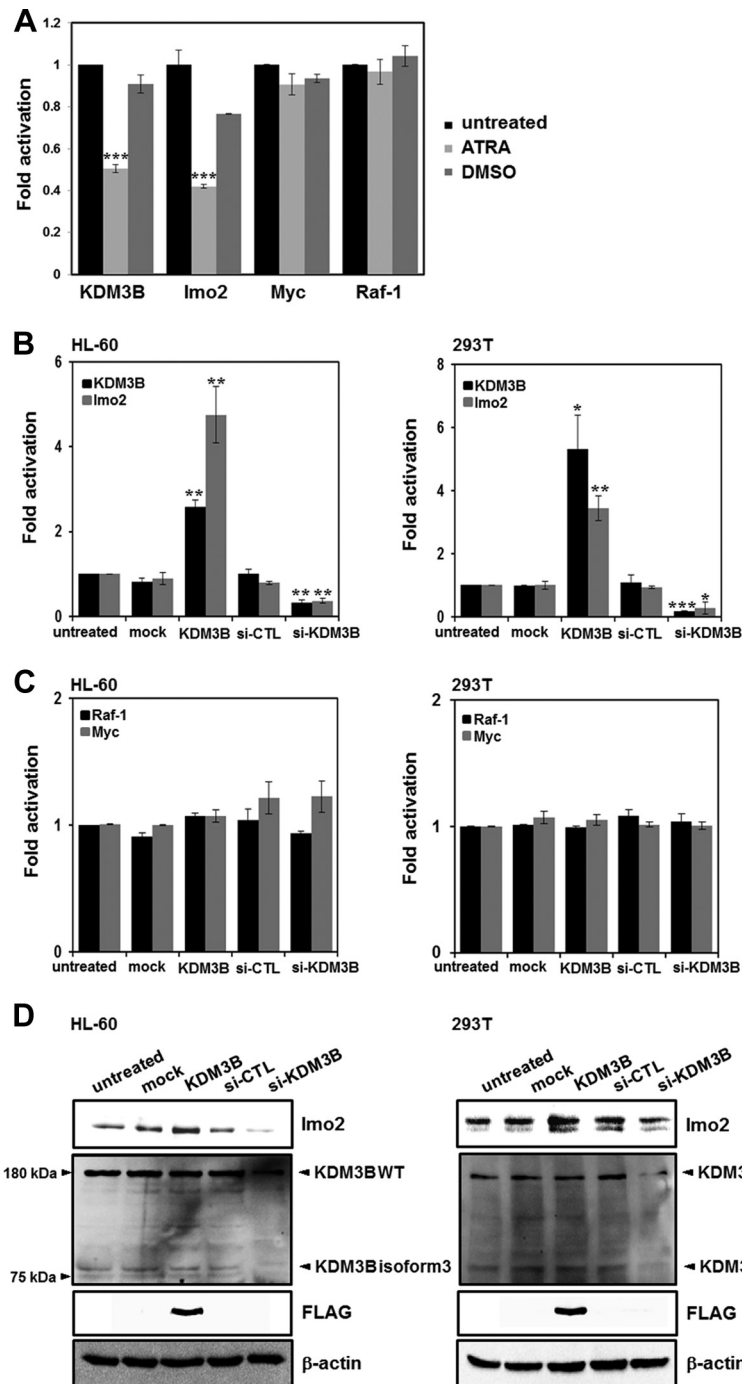


FIG 3 KDM3B activates the transcription of *lmo2*. (A) HL-60 cells were treated with ATRA or DMSO. After 48 h, real-time PCR was performed to compare the expression levels of the target genes. (B) The expression levels of *KDM3B* and *lmo2* in HL-60 and 293T cells with transiently overexpressed or knockdown *KDM3B* were detected via real-time PCR. The results are representative of at least three independent experiments (\pm SDs). *, $P < 0.05$; **, $P < 0.01$; ***, $P < 0.001$. (C) The expression levels of *Raf-1* and *Myc* in HL-60 or 293T cells with transiently overexpressed or knockdown *KDM3B* were detected via real-time PCR. The error bars represent $2^{-\Delta\Delta CT} \pm$ the SD of three independent experiments. (D) si-KDM3B-treated HL-60 and 293T cells were lysed and immunoblotted with anti-*lmo2* and anti-KDM3B antibodies. β -Actin was used as a loading control. (E) Flow chart showing the strategy used for microarray analysis and the *KDM3B* target gene identification process. (F) Identification of *KDM3B* target genes by hierarchical clustering; their expression changes in sh-CTL and sh-KDM3B knockdown stable cells were compared. Upregulated and downregulated gene clusters are represented by red and green, respectively. (G and H) Biological and molecular functional classification of *KDM3B* target genes. (I) The mRNA levels of *lmo2* and *KDM3B* in *KDM3B* knockdown stable cells were analyzed by real-time PCR. Results are shown as means \pm SDs; $n = 3$. ***, $P < 0.001$. (J) sh-CTL or sh-KDM3B knockdown stable cells were lysed and immunoblotted with anti-*lmo2* and anti-KDM3B antibodies. β -Actin was used as a loading control.

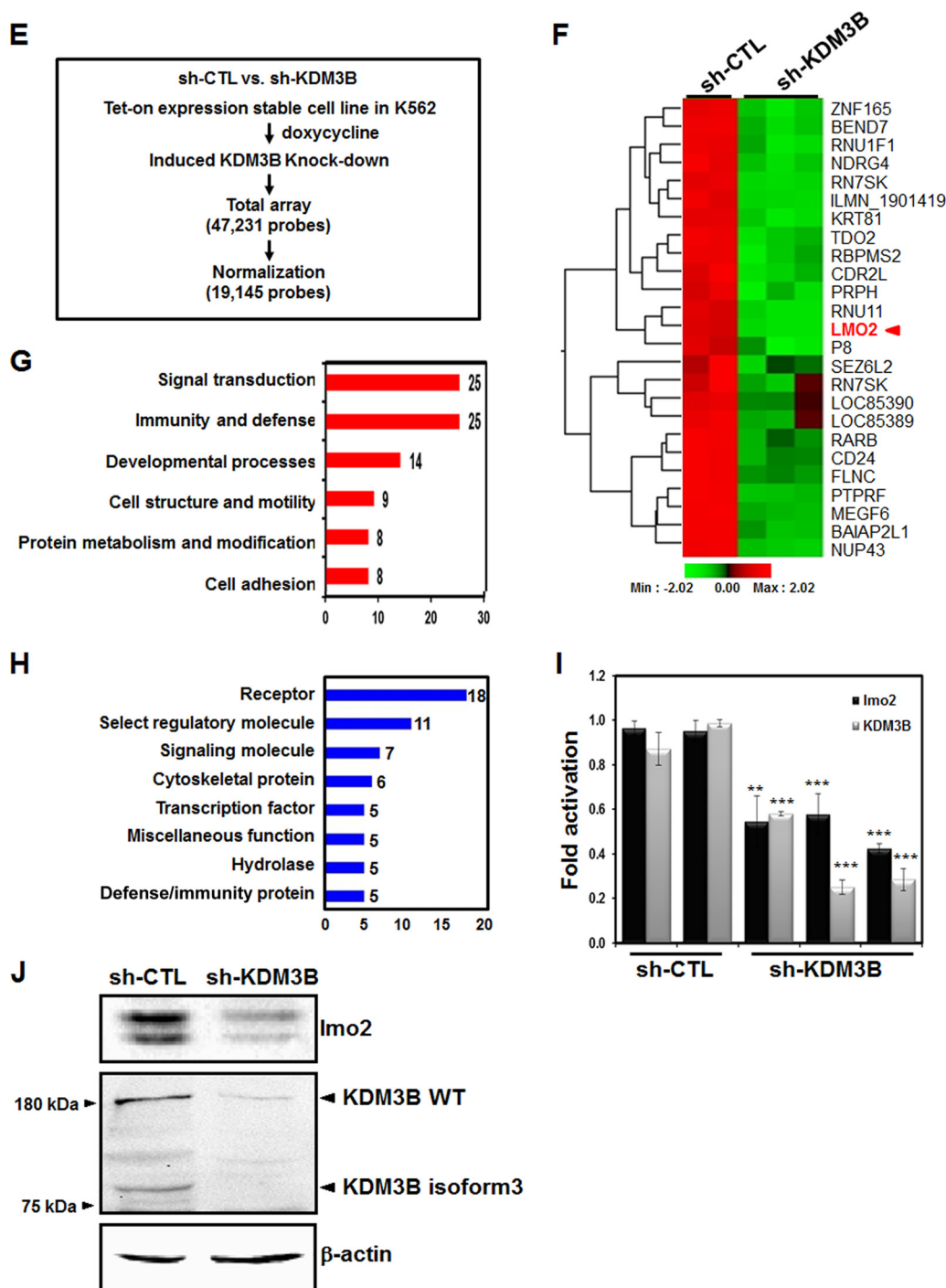


FIG 3 continued

pathways, such as *Raf-1* or *Myc*, was not altered (Fig. 3A). Next, we transfected HL-60 cells with KDM3B and determined the relative expression levels of selected oncogenes. In the real-time PCR analysis, the expression of *lmo2* was significantly increased upon KDM3B overexpression and decreased by si-KDM3B treatments (Fig. 3B). These results indicated that *lmo2* was a putative target gene regulated by KDM3B. In contrast, the expression of the other genes tested did not change (Fig. 3C). Protein expression levels

were also determined by immunoblot analysis. Knockdown of KDM3B using si-KDM3B clearly led to the downregulation of *lmo2* in HL-60 and 293T cells (Fig. 3D). Taken together, these results showed that KDM3B regulates transcription of *lmo2*.

To further investigate the effect of histone H3K9 demethylase KDM3B on leukemia, we performed global gene expression profiling with human erythroleukemic K562 cells stably expressing either control shRNA or KDM3B shRNA in a Tet-on inducible

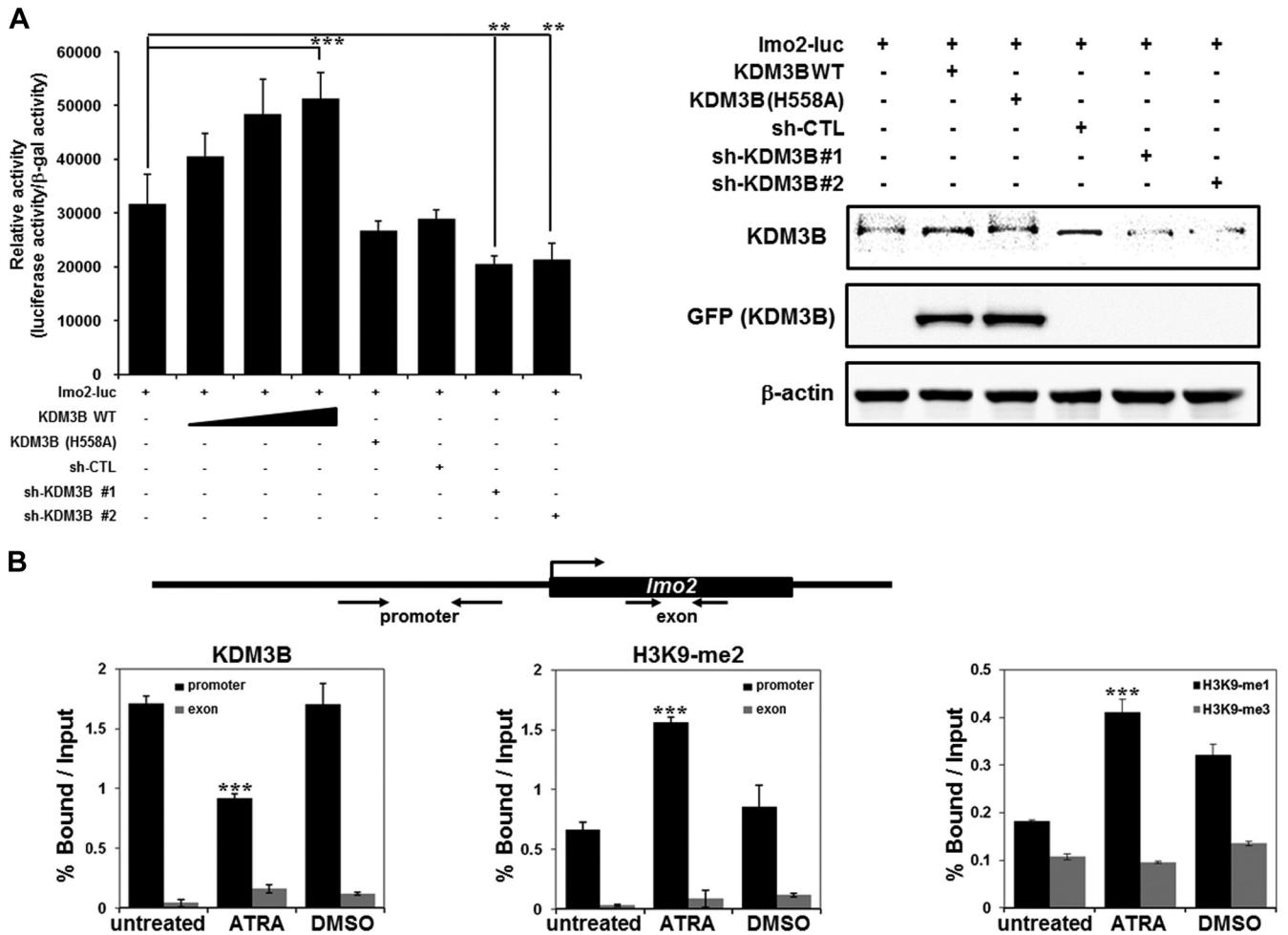


FIG 4 Recruitment of KDM3B to the *lmo2* promoter. (A) 293T cells were transfected with the pGL-*lmo2* promoter reporter and the indicated DNA constructs and sh-RNAs, and their cell extracts were assayed for luciferase activity. Luciferase activities were normalized to that of β -galactosidase. Each *P* value is the mean of five replicates from a single assay (left panel). The results are representative of at least three independent experiments (\pm SD). **, *P* < 0.01; ***, *P* < 0.001. Immunoblot analyses of the expression levels of endogenous and exogenous KDM3B in sh-KDM3B construct 1- or construct 2-transfected 293T cells were shown using anti-KDM3B and anti-GFP (right panel). (B) Schematic diagram of primer pairs in ChIP analysis (upper panel). Arrows indicate the primers used for real-time PCR amplification. ChIP analyses of the *lmo2* promoter in ATRA-treated HL-60 cells were conducted using anti-KDM3B (left panel), anti-H3K9-me2 (middle panel), and H3K9-me1/3 (right panel) and examined via real-time PCR. Results are the means \pm SDs; *n* = 3. ***, *P* < 0.001. (C to E) HL-60 (upper panel) or 293T (lower panel) cells were transfected with KDM3B and sh-KDM3Bs, and the ChIP assay and real-time PCR were performed using anti-KDM3B antibodies (C) to determine the methylation status of H3K9-me2 (D) and H3K9-me1/3 (E) in the *lmo2* promoter or exonic region. Results are the means \pm SDs; *n* = 3. *, *P* < 0.05; **, *P* < 0.01; ***, *P* < 0.001. (F and G) Schematic diagram of primer pairs in ChIP analysis (upper panel). Arrows indicate the primers for real-time PCR amplification. Transfected HL-60 cells with KDM3B and sh-KDM3Bs were analyzed by ChIP using anti-KDM3B antibodies. The immunoprecipitated DNA was amplified by PCR using the promoters of *Raf-1* (F) and *Myc* (G). The results are representative of at least three independent experiments (\pm SD). *, *P* < 0.05; **, *P* < 0.01; ***, *P* < 0.001.

expression system (Fig. 3E). Three RNA samples of sh-KDM3B and two control shRNAs were analyzed by a microarray (Fig. 3F). A total of 112 genes were either up- or downregulated by more than 1.5-fold when KDM3B was depleted (Fig. 3F and data available on request). In the biological function analysis using panther classification (<http://www.pantherdb.org>), these genes were shown to be associated with several major functional groups, including signal transduction, immunity and defense, developmental processes, cell structure and motility, and protein metabolism and modification (Fig. 3G). Furthermore, these differentially expressed genes were shown to be associated with other major groups, including receptors, regulatory molecules, signaling molecules, and transcription factors, by using molecular function

analysis (Fig. 3H). Notably, among the genes regulated by KDM3B, *lmo2* was found to be downregulated in the microarray analysis when KDM3B was depleted (Fig. 3F). Downregulation of *lmo2* in K562 cells stably expressing sh-KDM3B was further confirmed by real-time PCR and Western blot analysis (Fig. 3I and J).

KDM3B is recruited to the *lmo2* promoter and induces transcription. The LIM domain-only protein 2 (*lmo2*) gene is an important transcriptional regulator of erythropoiesis. Erythroid differentiation is completely blocked in both *lmo2*^{-/-} embryonic stem (ES) cells and wild-type ES cells transduced with α -*lmo2* antibodies (20, 30). To examine KDM3B-mediated transcriptional regulation of *lmo2*, we conducted a transfection assay using an *lmo2*-luc reporter system. Transcription of *lmo2* was enhanced

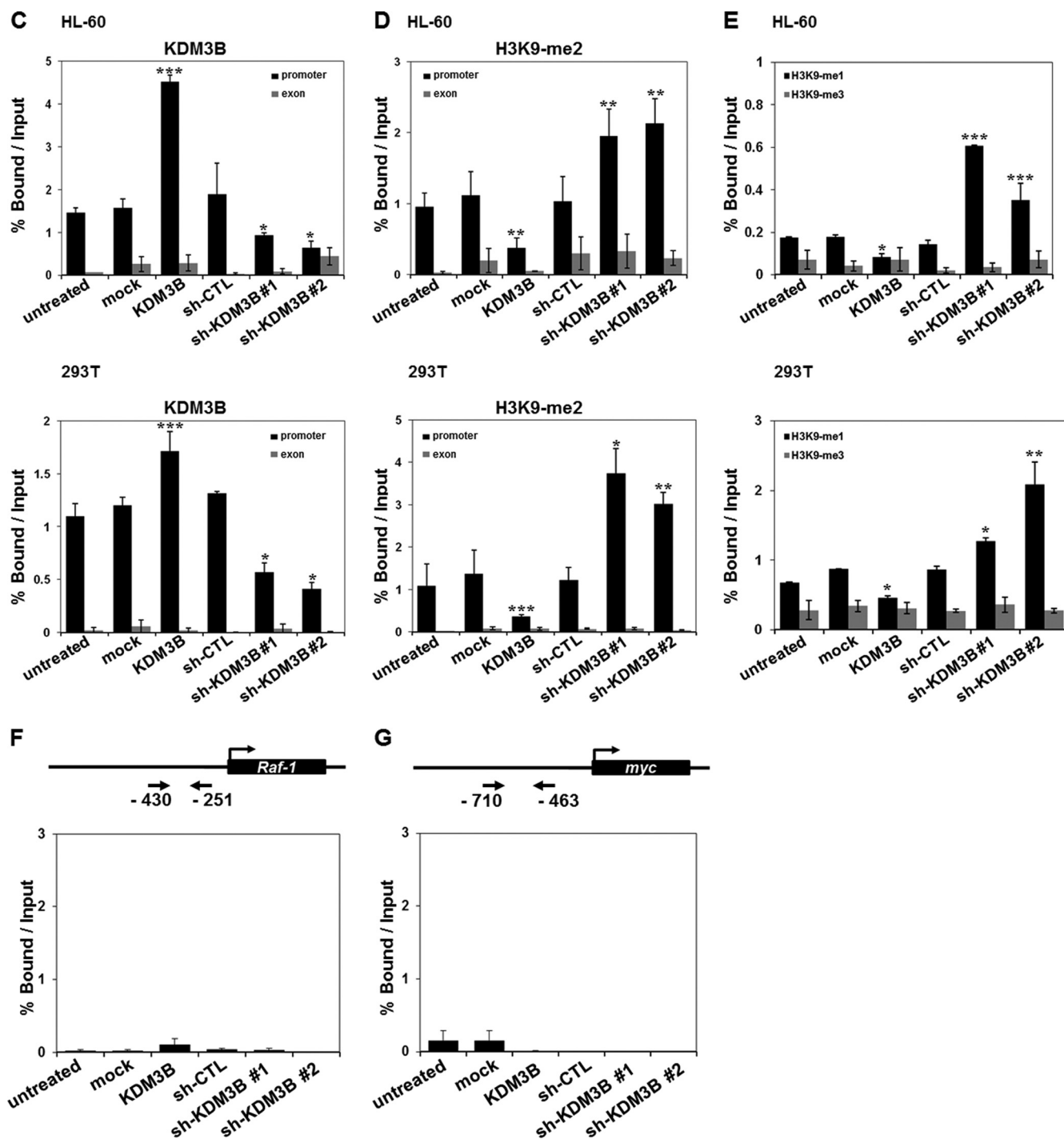


FIG 4 continued

by KDM3B cotransfection (Fig. 4A). Using the KDM3B H558A mutant, we verified that the transcriptional activation of *lmo2* by KDM3B was dependent on demethylase activity (Fig. 4A). Knockdown of KDM3B by two independent sh-KDM3Bs abolished transcriptional activation of *lmo2* mediated by KDM3B (Fig. 4A). Taken together, these findings indicate that KDM3B promotes leukemogenesis by activating the transcription of the hematopoietic oncogene *lmo2*.

To further investigate the mechanisms underlying *lmo2* transcriptional regulation via KDM3B, we performed ChIP analysis with real-time PCR using HL-60 cells. First, we observed decreased KDM3B recruitment as well as increased levels of H3K9-me2 and H3K9-me1 on the *lmo2* promoter in the presence of ATRA, results which were consistent with the initial results of the ChIP-chip analysis (Fig. 4B). However, H3K9-me3 levels did not change under the same conditions (Fig. 4B). Next, we evaluated

whether or not KDM3B can specifically target the *lmo2* promoter when overexpressed. Figures 4C and D show that KDM3B was highly recruited to the *lmo2* promoter, where it decreased H3K9-me2 methylation in both HL-60 (upper panel) and 293T (lower panel) cells. Knockdown of KDM3B by two different sh-KDM3Bs resulted in a significant reduction of KDM3B recruitment as well as an increase in H3K9-me1/2 methylation, but not H3K9-me3 methylation, on the *lmo2* promoter in both HL-60 and 293T cells (Fig. 4C to E). Specific recruitment of KDM3B to the *lmo2* promoter was validated by the absence of KDM3B recruitment to other oncogene promoters (*Raf-1* or *Myc*) when KDM3B was either overexpressed or knocked down in HL-60 cells (Fig. 4F and G). Collectively, these results suggest that KDM3B specifically targets the *lmo2* promoter and activates transcription through H3K9-me1/2 demethylase activity.

Functional connection between KDM3B and CBP. Having established that the histone demethylase KDM3B activates *lmo2* transcription by removing the epigenetic repression marker H3K9-me, it is reasonable to speculate that this transactivation process might be facilitated further by a KDM3B-interacting coactivator complex. To determine whether or not histone acetyltransferase (HAT) activity is involved in the KDM3B-mediated transcriptional activation of *lmo2*, transcriptional activation of *lmo2* was analyzed in the presence of HATs (CBP and PCAF) by using a luciferase reporter assay. Interestingly, KDM3B-mediated transactivation of *lmo2* increased in the presence of CBP, but this was not observed upon the addition of PCAF (Fig. 5A and B). Furthermore, a significant reduction in *lmo2* transcription by two independent sh-KDM3Bs, even in the presence of CBP, confirmed that KDM3B mediated coactivation with the HATs (Fig. 5A and B). This transactivation may have occurred due to the formation of a coactivator complex between KDM3B and CBP which may have induced *lmo2* expression. To test this hypothesis, immunoprecipitation assays between KDM3B and HATs were carried out. These assays clearly showed that CBP, but not PCAF, strongly interacted with KDM3B, forming an activator complex during *lmo2* transcription activation (Fig. 5C).

To verify whether CBP was recruited to the *lmo2* promoter via KDM3B, we performed ChIP analysis with real-time PCR using HL-60 cells. The CBP recruitment decreased on the *lmo2* promoter, as did the level of H3 acetylation in the ATRA-treated HL-60 cells (Fig. 5D and E). Also, CBP was highly recruited to the *lmo2* promoter when KDM3B was overexpressed, and the recruitment decreased upon KDM3B knockdown in HL-60 cells (Fig. 5F). Consistent with CBP recruitment, the H3 acetylation level either increased or decreased upon KDM3B overexpression or knockdown, respectively (Fig. 5G). These results suggest that KDM3B and the CBP HAT complex are both physically and functionally associated in transcriptional activation of *lmo2*.

KDM3B arrests HL-60 cell differentiation and is overexpressed in leukemia patient tissue samples. To investigate the potential role of KDM3B in leukemia cell differentiation, we examined the expression level of CD11b, a granulocytic differentiation marker, and CD14, a monocyte marker. The expression level of CD11b increased following ATRA treatment (63.7%) (Fig. 6A). Notably, KDM3B-transfected HL-60 cells clearly had a low expression level of the differentiation marker CD11b despite ATRA treatment (16.1%) (Fig. 6B). Interestingly, KDM3B knockdown by sh-KDM3B resulted in slight upregulation of CD11b expression (25.1%). Again, ATRA treatment in KDM3B knockdown

cells had significantly higher levels of the differentiation marker (65.6%) (Fig. 6C). However, the induction of CD14 did not change under the same conditions (Fig. 6A to C). The effect of KDM3B on HL-60 cell differentiation was further confirmed by morphological observation using Giemsa staining (Fig. 6D to F). Cells with more-prominent cytoplasmic granules and larger nuclei were dominant in undifferentiated conditions (Fig. 6D, untreated, and E, KDM3B+ATRA), whereas increasing fractions of cells showed more banded and segmented neutrophils in differentiating conditions (Fig. 6D to F, arrow-marked cells). These results suggest that KDM3B can prevent granulocytic differentiation of HL-60 cells by ATRA treatment. The contribution of *lmo2* regulation by KDM3B to leukemia cell differentiation was further investigated. As expected, the high expression level of CD11b induced by ATRA was downregulated by *lmo2* overexpression (Fig. 6G). Again, HL-60 cells showed increased levels of CD11b when *lmo2* was knocked down by si-*lmo2* in the absence of ATRA treatment compared to those of si-CTL (Fig. 6H). When *lmo2* was knocked down in the presence of ATRA, the CD11b expression level was further upregulated (72.6%). Finally, when we cotransfected KDM3B and si-*lmo2* in ATRA-treated HL-60 cells, we could clearly observe that knockdown of *lmo2* promotes HL-60 cell differentiation despite KDM3B overexpression (62.4%) (Fig. 6I). Those cells that underwent the differentiation process could be observed from undifferentiated cells by Giemsa staining and were marked with arrows (Fig. 6G to I, right panels). These results led us to the conclusion that regulation of *lmo2* by KDM3B is crucial for HL-60 cell differentiation.

Since KDM3B was identified in HL-60 cells and is highly expressed in tissues involved in hematopoiesis, we investigated whether or not KDM3B expression was dysregulated in leukemia patients by measuring the levels of KDM3B expression in bone marrow mononuclear cells. Total proteins were isolated from tissues recovered from eight healthy subjects and 30 patients with different types of leukemia. After isolating mononuclear cells, cellular phenotypes were examined by flow cytometry, and the characteristics of the cells are shown in Table S3 in the supplemental material. Using reverse transcription PCR (RT-PCR) analysis with primers specific for *KDM3B*, KDM3B was found to be overexpressed in selected leukemia patients compared to expression in normal cells (data not shown). A more-detailed analysis confirmed a higher level of KDM3B protein expression in ALL patients than in AML-type leukemia patients (Fig. 7A). To further correlate KDM3B and *lmo2* in ALL-type leukemia, we investigated whether the *lmo2* expression is also increased in ALL patient samples. Consistently, *lmo2* is more highly expressed in ALL-type patient samples than in AML-type patient and normal samples (Fig. 7B). These results suggest that KDM3B may play an important role via the regulation of *lmo2* in certain types of leukemogenesis.

DISCUSSION

In this study, we examined novel H3K9-me2 target genes during ATRA-mediated differentiation in HL-60 cells using ChIP-chip analysis. Among the target genes, the histone demethylase KDM3B was shown to be downregulated during leukemia cell differentiation via H3K9-me2 modification at the promoter.

Histone methylation-demethylation dynamics have been extensively studied and are recognized as important markers for a variety of biological processes, including transcription regulation.

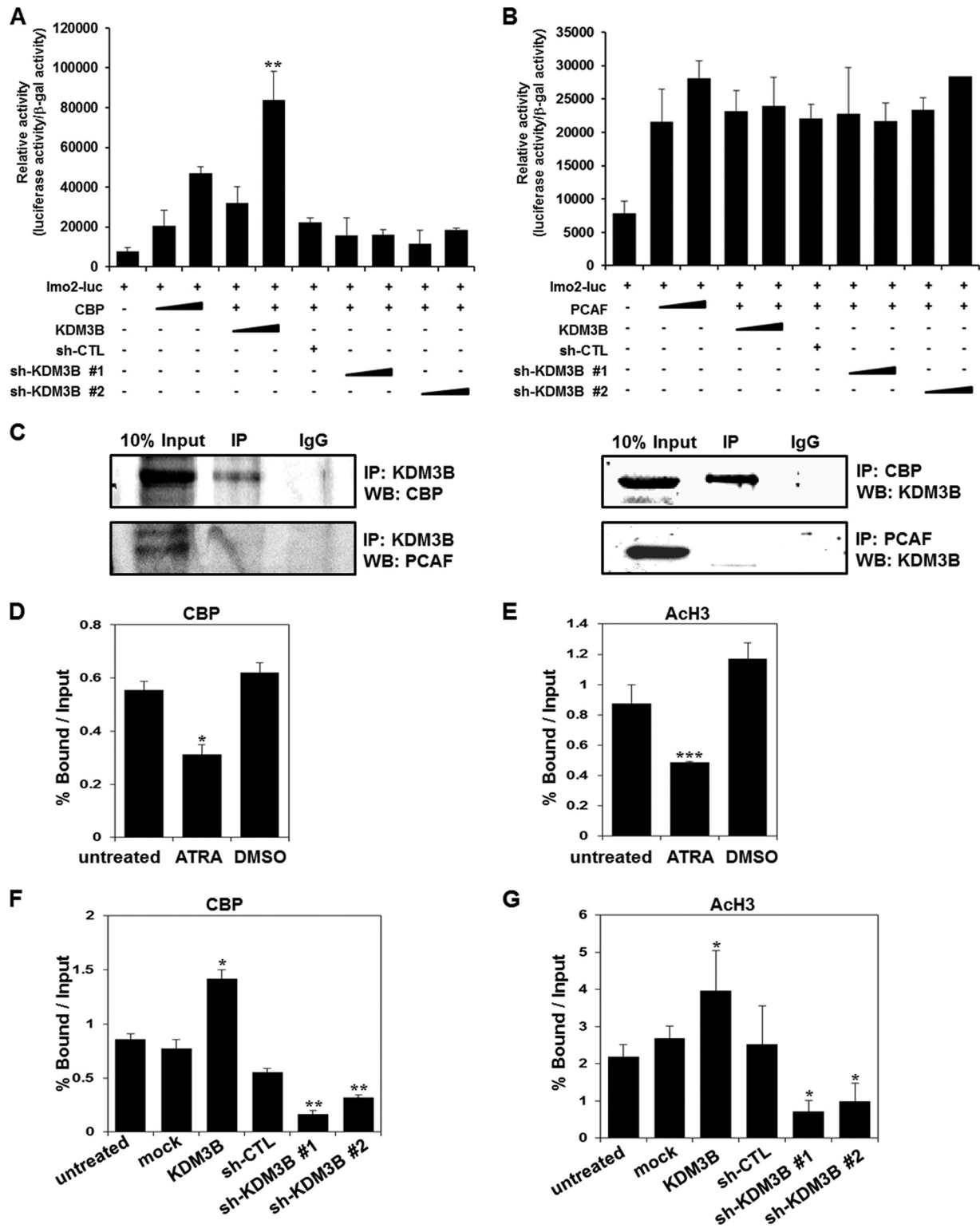


FIG 5 KDM3B regulates *lmo2* transcription via interaction with CBP. (A and B) 293T cells were transiently transfected with the *lmo2* promoter reporter plasmid along with KDM3B, sh-KDM3Bs, and CBP (A) or PCAF constructs (B). The results are representative of at least five independent experiments (\pm SD). **, $P < 0.01$ compared with the untreated control. (C) 293T cells were transfected with pcDNA6-KDM3B, Gal4-CBP, and FLAG-PCAF. Immunoprecipitation was performed using anti-KDM3B, anti-CBP, anti-PCAF, or anti-immunoglobulin G (IgG) antibodies. A Western blot analysis was then conducted using anti-CBP, anti-PCAF, or anti-KDM3B antibodies. (D and E) ChIP analyses of the *lmo2* promoter in ATRA-treated HL-60 cells were conducted using anti-CBP (D) or anti-ACh3 (E) antibodies. The results are expressed as means \pm SDs; $n = 3$. **, $P < 0.01$; ***, $P < 0.001$. (F and G) HL-60 cells were transfected with KDM3B and sh-KDM3Bs, and the ChIP assay and real-time PCR were performed in the *lmo2* promoter using anti-CBP (F) and anti-ACh3 (G) antibodies. The results are the means \pm SDs; $n = 3$. *, $P < 0.05$; **, $P < 0.01$; ***, $P < 0.001$.

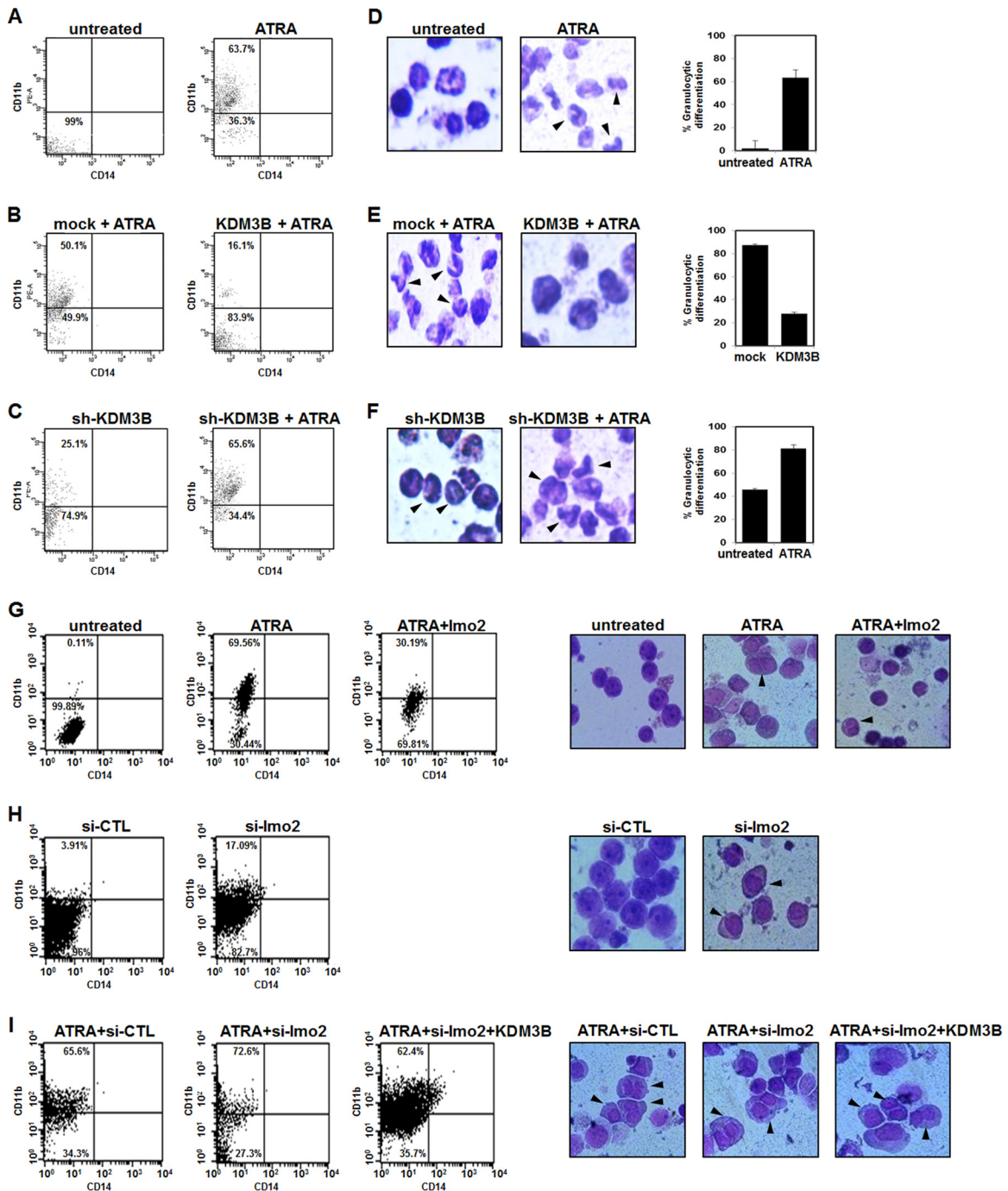


FIG 6 KDM3B induces leukemic transformation. (A to C) Granulocytic differentiation was analyzed by FACS 72 h after ATRA treatment. Percentages of CD11b-positive cells in HL-60 cells after transfection with KDM3B or sh-KDM3B as follows: ATRA (A), KDM3B with ATRA (B), or sh-KDM3B with ATRA (C). (D to F) Morphological analysis and graphic representation of the morphological data of HL-60 cells 72 h after transfection with KDM3B or sh-KDM3B as follows: ATRA (D), KDM3B with ATRA (E), or sh-KDM3B with ATRA (F). (G and H) CD11b/CD14 expression following transfection with lmo2 or si-lmo2 by FACS: panel G, lmo2 with ATRA; panel H, si-lmo2. (I) HL-60 cells were treated with ATRA (1 μ M) during 72 h after transfection with si-CTL, si-lmo2, or a combination of si-lmo2 and KDM3B (left panels). Cell morphology of ATRA-treated HL-60 cells analyzed by Giemsa staining after transfection with lmo2, si-lmo2, or KDM3B (right panels).

KDM3B, containing the JmjC domain, is a member of the KDM3 family that includes KDM3A (testis-specific gene A [TSGA]) and KDM3C (hormone receptor-interacting protein 8 [TRIP8]), which show H3K9-me1/2 demethylase activity in human and

mice, respectively (15, 31). Initially, the gene was identified as 5q nuclear coactivator (5qNCA) and thought to have possible functions in myelodysplastic syndrome (MDS) and acute myeloid leukemia (AML) (14). KDM3A was also found to play a role in mouse

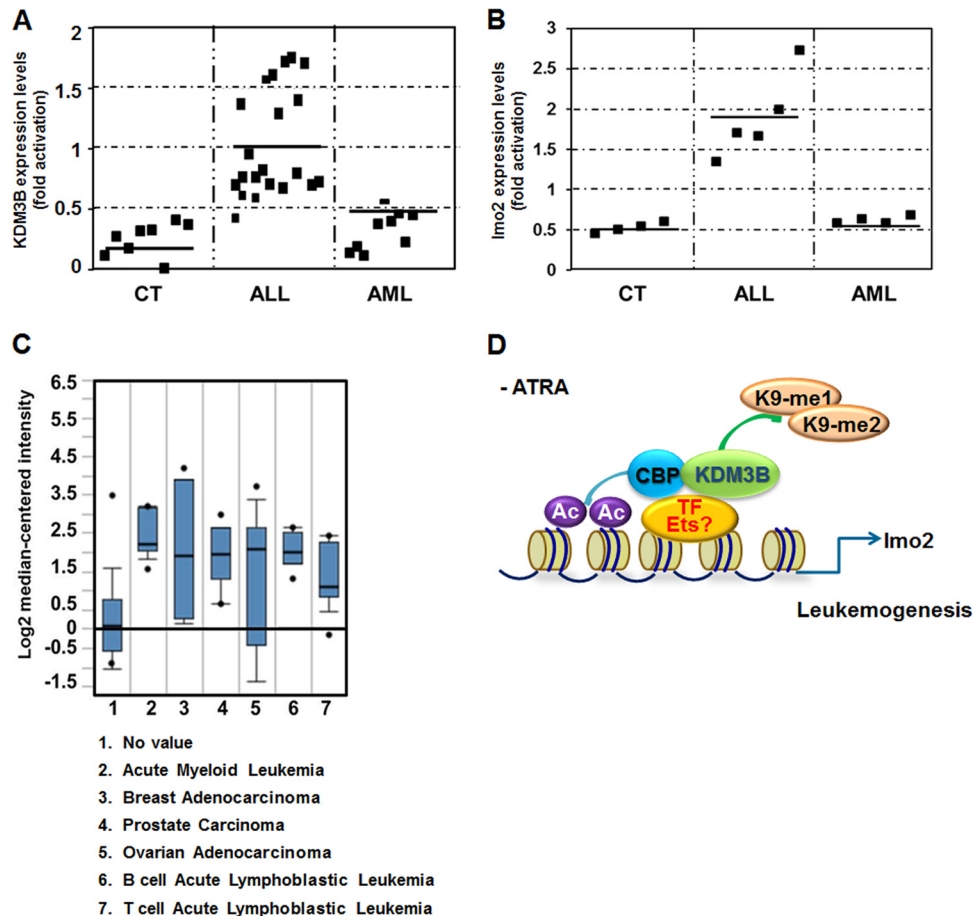


FIG 7 KDM3B was overexpressed in ALL-type leukemia patient samples. (A and B) Each individual spot of KDM3B (A) and *lmo2* (B) expression in total cells extracted from blood cells of healthy individuals and leukemia patients. Western blots were analyzed quantitatively in the spots by the Image J program. The data are averages of two independent experiments. (C) Boxplot of KDM3B expression in normal and cancerous tissues. 1, no value; 2, acute myeloid leukemia, $P = 4.98E-5$; 3, breast adenocarcinoma, $P = 0.3E-5$; 4, prostate carcinoma, $P = 4.3E-4$; 5, ovarian adenocarcinoma, $P = 2.65E-4$; 6, B cell acute lymphoblastic leukemia, $P = 1.83E-3$; 7, T cell acute lymphoblastic leukemia, $P = 4.2E-4$. All P values are calculated in comparison with column 1. (D) Model for regulation of *lmo2* transcription by KDM3B in leukemogenesis.

spermatogenesis and obesity (21, 26). KDM3C was identified as an androgen receptor coactivator that regulates mouse steroidogenesis through H3K9 demethylation at the *p450c17* promoter (15, 21). Our current study revealed that KDM3B is an H3K9-me1/2 demethylase that displays specific activity *in vivo* and *in vitro* in leukemogenesis and acts as a transcriptional coactivator.

Using both global gene expression profiling and searching for individual target genes, *lmo2* was identified as a possible target gene of KDM3B. *lmo2* is an important transcriptional regulator of erythropoiesis and has been reported to inhibit erythroid differentiation (18, 28). Studies on the epigenetic regulation of *lmo2* transcription recently found that JAK2 phosphorylates H3Y41 and that HP1 α exclusion results in *lmo2* activation in leukemia cells (7). We demonstrated that KDM3B activated the expression of *lmo2* through increased H3K9-me1/2 demethylase activity at the *lmo2* promoter. Identifying upstream regulators of JAK2 and its impact on *lmo2* in leukemia will be the topic of further studies.

Reports have shown that histone demethylases in multicomplexes was associated with other chromatin remodelers. For example, the H3K27 demethylase UTX, in association with the H3K4 HMTase MLL2, leads to the activation of target genes, such

as *Hox* (4, 19). Interestingly, we found that KDM3B interacted with CBP but not PCAF, and this association further enhanced transcriptional activation of *lmo2*. Among the different HATs, CBP regulates various hematopoietic transcription factors and is involved in hematopoietic development and associated diseases (2). Another report suggested that different epigenetic mechanisms, such as histone deacetylase (NuRD) and demethylase (LSD1), interdependently regulate the transcription of key target genes involved in breast cancer (29). Therefore, it is not surprising that simultaneous histone demethylation of H3K9 by KDM3B and acetylation of H3/H4 by CBP activate target gene expression. Identification of multiple effectors forming a complex via direct binding during chromatin remodeling and subsequent transcriptional regulation of tissue-specific gene expression should allow for the further characterization of coactivation complexes, including those containing KDM3B and CBP, as well as their impact on specific gene expression during leukemogenesis.

Recent reports have suggested that histone demethylases, such as KDM4C (GASC1), KDM5B (Plu-1), and KDM8, play important roles in carcinogenesis (5, 10, 12). KDM4C is believed to cause HP1 delocalization through its H3K9-me2/3

demethylase activity and reverse oncogene silencing (5). Involvement of histone demethylases in carcinogenesis, particularly for leukemia, has been suggested by a number of recent studies. LSD1 is known to prevent H3K4 methylation mediated by MLL. Inhibition of LSD1 perturbs the differentiation of cells with a hematopoietic lineage (24). The hematopoietic transcription factor TAL1 is also negatively regulated by LSD1 (13). Furthermore, depletion of the H3K36 demethylase KDM2b/JHDM1b significantly inhibits leukemogenesis mediated by Hoxa9/Meis1 (11). To investigate whether or not KDM3B can induce leukemic proliferation, we performed a cell differentiation assay. Initially, the expression profile of the differentiation marker CD11b was found to be either downregulated or upregulated in overexpressed or knockdown KDM3B cells, respectively. These results indicate that KDM3B may modulate the differentiation of the leukemia cell line HL-60. Furthermore, lmo2 overexpression has been shown to offset ATRA-mediated differentiation, and knockdown of lmo2 promotes differentiation even though KDM3B was overexpressed. Interestingly, overexpression of KDM3B was observed in ALL-type leukemia patients but not AML-type leukemia patients. The reason for this difference is not clear. However, the human *KDM3B* gene is located at 5q31, and this region is frequently deleted in AML and myelodysplasia (14). It has been previously reported that there is a clear difference in gene expression patterns between AML and ALL. The study suggested that human leukemia subtype class discovery and class prediction between AML and ALL can be made based on gene expression profiles by DNA microarray (9). The significance of this specificity probably needs further investigation. The finding that lmo2 was also overexpressed in ALL-type patients further links the role of KDM3B in regulation of lmo2 expression to this particular type of leukemia.

To obtain further evidence of KDM3B involvement in leukemogenesis, we searched the Oncomine database to compare the expression levels of KDM3B in different types of leukemia to those in normal tissues. KDM3B was found to be upregulated in various types of leukemia compared to its expression in normal tissues (Fig. 7C). These data strongly suggest that KDM3B plays a role in leukemia, possibly through H3K9-me1/2 demethylation.

In summary, we have provided *in vitro* and *in vivo* evidence for the possible role of the H3K9-me1/2 demethylase KDM3B in leukemia through activation of the leukemic oncogene *lmo2* (Fig. 7D). Our model suggests the existence of a transcription factor for lmo2 associating with KDM3B and CBP and transactivating *lmo2* expression in the absence of ATRA (Fig. 7D). In fact, regulation of *lmo2* transcription by Ets transcription factors and Ets-1 transactivation potentiation by CBP/p300 have been reported (18, 32). On the other hand, ATRA-dependent RAR/RXR may activate transcription of an unidentified KDM3B repressor and this repressor downregulates transcription of KDM3B (after ATRA treatment). We examined the epigenetic regulation of target gene transcription by KDM3B during ATRA-induced leukemia cell differentiation. Taken together, our results demonstrate the role of H3K9 demethylase KDM3B in the proliferation of leukemic cells through its epigenetic regulatory activity.

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