MOZ Forms an Autoregulatory Feedback Loop with miR-223 in AML and Monocyte/Macrophage Development

HIGHLIGHTS
- Low MOZ is associated with poor prognosis in patients with AML.
- Silencing of MOZ makes AML cells exhibit "stem-like" characters.
- MOZ promotes monocyte-to-macrophage development and M1 polarization.
- MOZ is a direct target of miR-223 and functions as a repressor of miR-223.

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MOZ Forms an Autoregulatory Feedback Loop with miR-223 in AML and Monocyte/Macrophage Development

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SUMMARY
Monocytic leukemia zinc-finger protein (MOZ) has been found to form fusion proteins with many regulators in acute myeloid leukemia (AML). However, the molecular functions and underlying mechanism of MOZ in AML is not well understood. Here, clinical MOZ expression analysis combined with data integration from the TCGA and GEO databases indicated that a low level of MOZ was associated with poor prognosis. MOZ knockdown inhibited monocyte differentiation and increased resistance to chemotherapeutic drug-induced apoptosis in THP-1 or U937 cells. In addition, we found that genetic silencing of MOZ suppressed AP-1 and AKT activity in the context of lipopolysaccharide stimulation, resulting in diminished M1 activation of macrophages. We further showed that MOZ was a validated target of miR-223 and functioned as a repressor of miR-223 expression. Our study indicates that a molecular network involving MOZ and miR-223 contributes to the monocyte differentiation and polarization program, which is deregulated in AML.

INTRODUCTION
Monocytes that originate from hematopoietic stem cells in the bone marrow have been considered the systemic reservoir of myeloid precursors for the renewal of macrophages and dendritic cells (DCs) (Iwasaki and Akashi, 2007). Numerous master regulators, including PU.1, C/EBPα, FLT3, and miR-223, have been reported to participate in monocyte differentiation (Friedman, 2007; Gilicze et al., 2014). However, somatic mutation or genetic variation of these master regulators in monocytes often causes hyperplasia and blocks cellular differentiation, resulting in the clinical entity acute monocytic leukemia (AMoL) (Palmisano et al., 2008; Renneville et al., 2008). As immune effector cells, macrophages are directly derived from monocytes under inflammatory conditions (Ginhoux and Jung, 2014). In response to different stimuli, macrophages can be induced to differentiate along two main functional pathways: classical (M1) activation induced by interferon-γ or Toll-like receptor (TLR) agonists and alternative (M2) activation induced by Th2 cytokines such as interleukin (IL)-4 or IL-13 (Italiani and Boraschi, 2014). M1 macrophages produce high levels of pro-inflammatory factors and inducible NO synthase (iNOS), which are critical for eradicating bacterial, viral, and fungal infections. M2 macrophages are characterized by the high expression of markers of alternative activation, such as arginase-1 (Arg-1) and IL-10, which participate in the response to parasitic infection, tissue remodeling, and tumor progression (Wang et al., 2014). At the molecular level, many transcription factors (TFs), such as nuclear factor (NF)-kB, AP-1, and PU.1, participate in TLR-induced M1 activation, whereas STAT3 and C/EBPβ are involved in the polarization of macrophages to the M2 phenotype (Lawrence and Natoli, 2011; Murray and Wynn, 2011; Mosser and Edwards, 2008). The epigenetic modifiers JMJD3 and HDACs have also been reported to participate in M2 activation (Ivashkiv, 2013).

Monocytic leukemia zinc-finger protein (MOZ) is a founding member of the MYST family of lysine acetyltransferases, together with Ybf2/Sas3, Sas2, and TIP60 (Snyder and Glass, 2000). MOZ has intrinsic histone acetyltransferase (HAT) activity, increasing the acetylation of H3K14 and H3K9, leading to chromatin remodeling (Carlson and Glass, 2014; Champagne et al., 2001). MOZ appears to function as a cofactor for AML1 (RUNX1), PU.1, and p53-mediated gene expression (Kitabayashi et al., 2001; Rokudai et al., 2013; Katsumoto et al., 2004). Recurrent translocations within the MOZ gene are associated with AMoL (Troke et al., 2006). Fusion proteins such as MOZ-TIF2 and MOZ-NCOA3 retain the N-terminal portion of MOZ and show aberrant functionality, which affects histone modification and gene regulation (Deguchi et al., 2003; Esteyries et al., 2008). In line with the evidence linking MOZ to leukemia, some...
studies have highlighted the essential role of MOZ in the development and maintenance of hematopoietic stem cells (Katsumoto et al., 2006; Perez-Campo et al., 2013; Sheikh et al., 2016). Owing to the lack of a mouse model (the MOZ−/− genotype is embryonic lethal), the molecular functions of MOZ involved in monocyte differentiation and macrophage activation are poorly understood.

MicroRNA (miRNAs) are short, endogenous, single-stranded RNA molecules that regulate gene expression at the posttranscriptional level (Bartel, 2004). Recent studies have shown that a range of miRNAs have critical functions in the regulation of inflammatory responses in macrophages and monocytes (Lu and Liston, 2009; Gilicze et al., 2014). miR-223 was in the first cadre of miRNAs discovered to be highly expressed in myeloid cells. The expression of miR-223 is regulated by a combination of factors, including PU.1, E2F1, GATA1, C/EBPα, and C/EBPβ (Fukao et al., 2007; Pulikkan et al., 2010; Eyholzer et al., 2010; Zhou et al., 2015). miR-223 has been proved to play an important role in granulocyte differentiation by targeting E2F1, and deregulated miR-223 is associated with acute myeloid leukemia (AML) progression (Pulikkan et al., 2010). Before IKKα and STAT3 were confirmed as validated targets of miR-223 in recent studies, it was believed that miR-223−/− macrophages have a stronger response to lipopolysaccharide (LPS) stimulation.

Our initial observation of the significantly reduced MOZ expression in patients with AMoL prompted us to investigate the potential effect of MOZ on AMoL pathogenesis. In this study, knockdown of MOZ in THP-1 cells blocked phorbol myristate acetate, a specific activator of Protein Kinase C (PKA)-induced differentiation and conferred apoptotic resistance to chemotherapeutic-drug-induced cell death. Loss of MOZ diminished TLR-induced M1 activation of macrophages. Our results also demonstrate that MOZ is a validated target and transcriptional repressor of miR-223. Thus our study shows that monocyte differentiation and macrophage activation are regulated by the MOZ-miR-223 network, in which MOZ functions as a key regulator of differentiation and polarization, interlinked with miR-223 in a mutual negative feedback loop.

RESULTS
Low Expression of MOZ Is Associated with Poor Prognosis in AML Cases

The pathogenesis of AML involves the abnormal proliferation and blocked differentiation of a clonal population of myeloid stem cells. Various biomarkers for AML have been identified, such as FLT3-ITD, NPM1 mutants, and C/EBPα mutants. Based on previous reports, MOZ has been found to be fused to other moderators (NCOA3, TFII, p300, etc.) in AMoL caused by chromosome translocation (Esteyries et al., 2008; Carapeti et al., 1997; Chaffanet et al., 2000). However, the specific role of MOZ in AML pathogenesis is not well understood. Here, we applied an integrated approach and analyzed multidimensional data from AML cases in the TCGA and GEO databases to investigate the relationship between MOZ and AML. As shown in Figure 1A, we found low expression of MOZ in FLT3-TKD, FLT3-ITD, and NPM1 mutant groups, as well as high expression in a C/EBPα mutant group. Given that FLT3-TKD and FLT3-ITD mutant patients were associated with poor outcomes in AML, whereas improved prognosis was noted in C/EBPα mutant patients, our results suggested that low expression of MOZ was associated with aggravated AML progression. In addition, we performed a correlation analysis between MOZ and well-defined AML biomarkers in AML cases in the TCGA cohort. As shown in Figure 1B, we found that MOZ was inversely associated with various markers. In addition, we compared the MOZ expression among different subgroups of AML based on the French–American–British (FAB) classification system. Interestingly, we found that the expression of MOZ was lowest in the M5 group in TCGA or GEO cohorts, as presented in Figure 1C. To determine whether the monocyte proportion in leukemia contributed to the observed low expression of MOZ in the M5 group, we compared the log2-transformed expression value of MOZ in different AML cases grouped by the proportion of monocytes. Indeed, as shown in Figures 1D and 1A, a high percentage of monocytes was associated with low MOZ expression. To exclude the possibility of intrinsic expression differences, we compared the expression of MOZ in monocytes and other lymphocytes. No significant difference was observed between CD14+ monocytes and CD14+ lymphocytes (derived from peripheral blood mononuclear cells), as shown in Figure 1D, suggesting that MOZ was not heterogeneously expressed in diverse cell groups, at least in hematopoietic-derived cell subgroups. Furthermore, M5 patients younger than 65 years with low expression of MOZ in the TCGA cohorts with normal karyotypes showed worsened overall survival, whereas GEO cohorts showed the same trend (Figure 1F). Collectively, integrated analysis of the TCGA and GEO cohorts suggested that low MOZ was strongly associated with poor clinical outcomes in AMoL.
MOZ Knockdown Aggravates AMoL Progression

To validate the correlation between AMoL development and MOZ expression, we examined the MOZ transcription level in peripheral blood samples from patients with AMoL and healthy volunteers. By using unique primers that were designed to abandon the samples containing MOZ fusion fragments caused by chromosome translocation, we found that MOZ expression was significantly reduced in AMoL samples, as presented in Figure 2A. Compared with the level in purified monocytes from healthy individuals, a lower level of MOZ was observed in THP-1 cells, which are broadly used as a model for monocyte development and pathogenesis in leukemia research (Figure 2A). As altered MOZ expression was observed in AMoL, we sought to determine the expression of MOZ during normal monocyte differentiation. As shown in Figures 2B and 2D, sustaining increased MOZ expression was observed during cytokine-induced primary monocyte development. Similar results were obtained in PMA-induced or 1,25(OH)2-VitD3 (VD3)-induced THP-1 transformation (Figures 2C and S1A). In addition, MOZ was also upregulated in PMA- and VD3-induced U937, which is another monocytic AML cell line (Figure S1B). These results suggest that there is a causal connection between MOZ expression and monocyte fate. To address this hypothesis, we established a stable transfected THP-1 or U937 cell line expressing short hairpin RNA against MOZ transcripts (designated as shMOZ), and the efficiency of knockdown was confirmed (Figures 2E and S1D). In our experiments, we found that the stem cell maker CD34 was up-regulated in shMOZ cells, whereas loss of MOZ reduced CD38 or CD11b expression when compared with control cells (Figures 2F and S1C). In addition, MOZ knockdown arrested the cell cycle in THP-1 cells as evidenced by more cells in the G0 phase (Figure 2G). Similar results were observed in U937/shMOZ cells, which exhibited decreased proliferation activity (Figure S1E). Vimentin and E-cadherin are well-characterized differentiation-related markers (Mor-Vaknin et al., 2003; Van den Bossche et al., 2015). Here, we found that the level of vimentin, which is down-regulated by induction with PMA, was higher in shMOZ than in control cells, whereas PMA-induced up-regulation of E-cadherin was significantly lower in shMOZ cells. In addition, shMOZ cells exhibited

Figure 1. Low Expression of MOZ Is Associated with Poor Prognosis in TCGA and GEO Cohorts

(A) Statistical analysis of MOZ expression in somatic mutant and normal groups in the GEO or TCGA cohorts.
(B) Correlation analysis between MOZ and AML-related biomarker expression in the TCGA database using SPSS.
(C) MOZ expression in different subgroups of AML based on the FAB classification system.
(D) Comparison of MOZ expression in definite groups with differential proportions of monocytes.
(E) Isolated CD14+ monocytes and eluted lymphocytes from human peripheral blood mononuclear cells were used for determining the mRNA level of MOZ by quantitative PCR.
(F) Survival analysis of MOZ in patients with AML (M5 subtype younger than 65 years) in TCGA (hazard ratio [HR] = 4.01) or GEO cohorts (HR = 1.37) was performed using Kaplan-Meier methods. *p < 0.05; **p < 0.01; ***p < 0.001; NS, not significant.
Figure 2. Knockdown of MOZ Promotes AMoL Development

(A) Sorted monocytes from peripheral blood of patients with AML or healthy individuals (left) or immortal THP-1 cells (right) were lysed for quantitative PCR (qPCR), using MOZ-specific primers.

(B–D) Monocytes from mouse bone marrow or human peripheral blood and THP-1 cells were treated with M-CSF (20 ng/mL), GM-CSF (100 ng/mL), or PMA (10 μg/mL) for the indicated time periods. qPCR was performed to determine MOZ expression, normalized to GAPDH (B and C). Representative western blot (D) of MOZ is shown.

(E) The efficiency of MOZ knockdown in THP-1 cells is shown.

(F) shMOZ stable-transfected THP-1 cells and control cells were labeled with anti-CD34 or anti-CD38 antibodies conjugated with fluorescein isothiocyanate or phycoerythrin (PE) and analyzed using flow cytometry.

(G) THP-1 shMOZ or control cells were cultured for 24 hr, fixed, and permeabilized with ethanol (100%) overnight. Sequentially, the cells were stained with propidium iodide (PI) at the indicated concentration and examined for cell division using fluorescence-activated cell sorting (FACS).
an impaired differentiation progression, based on lower PMA-induced adherence and limited production of reactive oxygen species (ROS) triggered by LPS stimulation compared with that in control cells (Figures 2I and S1I). Furthermore, the loss of MOZ conferred apoptosis resistance to THP-1 cells or U937 against cisplatin-induced cell death (Figures 2J, S1F, and S1G, statistical results shown on the right). Besides, Wright-Giemsa staining showed that higher nuclear-cytoplasmic ratio was present in MOZ knockdown cell (Figure S1H). Taken together, these data confirm the critical role of MOZ in monocyte development and show that loss of MOZ blocks normal differentiation and facilitates AMoL pathogenesis.

Furthermore, we also used specific inhibitor (MOZ-IN-3) targeting against MOZ to evaluate the role of MOZ in U937. As shown in Figure S2B, U937 treated with MOZ-IN-3 for 15 days had higher nuclear-cytoplasmic ratio induced with PMA or VD3 when compared with non-treated cells. Also, enhanced anti-apoptosis activity to chemotherapeutic-drug-induced cell death was observed in U937 treated with MOZ-IN-3 (Figures S2C and S2D). In line with the observation in MOZ knockdown U937, these results confirmed that MOZ activity was critical for monocyte functions.

Dysregulated MOZ Has Strong Connection with Mutated Myeloid-Related Regulators in AML

To uncover the reason why MOZ is expressed at a low level in AML, especially in AMoL, we screened the sequence 3 kb upstream of the proximal TSS (transcription start site) of the human MOZ gene for regulatory elements and corresponding factors using various software available online as detailed in Figure 3A. Subsequently, the predicted TFs with high scores were organized and ranked for GO analysis. As shown in Figure 3B, the identified TFs were strongly associated with AML pathogenesis. Based on the knowledge of somatic mutation in some master genes contributing to tumorigenesis, we analyzed the mutant frequency of predicted TFs in various cancer cases in the TCGA cohort. As shown in Figure 3C, we found that most of these factors possess > 10% mutant rate among the 500 most mutant cases in the TCGA database. Furthermore, gene co-expression analysis performed using cytoscape software (GeneMINIA module) indicated that MOZ expression was strongly connected with these chosen TFs, especially SP1, SPI-1, and EST-1 (Figure 3D). In addition, the co-expression network comprised by these TFs and the 10 factors with the highest mutation rate in AML indicated that they shared extensive functional properties (Figure 3E). Thus based on the above bioinformatic analysis, it is reasonable to assume that impaired MOZ expression in AMoL might be attributable to the abnormal activity of AML-related regulators caused by somatic mutation or deregulation mediated by other modulators.

Loss of MOZ Diminishes the M1 Activation of Macrophage

The above results show that MOZ plays an important role in determining monocyte fate (normal differentiation versus AML pathogenesis). As immune effector cells, macrophages are derived directly from monocytes under normal or inflammatory conditions (Varol et al., 2015). To determine whether MOZ takes part in macrophage functions, we measured the expression of MOZ in activated macrophages induced by TLR agonists. As shown in Figures 4A and 4B, MOZ was clearly up-regulated after ligand stimulation. It is well accepted that TLR ligands (LPS, poly I:C, or PAM3CK4) induce M1 polarization, whereas the cytokines IL-4 and IL-13 polarize cells to the M2 phenotype, initiating the tolerance reaction (Wang et al., 2014). Here, IL-4 induced the down-regulation of MOZ expression but elevated arginase expression in knockdown cells, indicating that the loss of MOZ was favorable for M2 polarization (Figures 4C and S1A). To further explore the functions of MOZ in macrophages, we generated a puromycin-resistant RAW264.7 cell line that was stably transfected with the expression cassette of a small interfering RNA targeting MOZ transcripts and pac gene coding for an enzyme that degrades puromycin (Figure 4D). MOZ knockdown macrophages (designed as shMOZ) showed lower production of pro-inflammatory factors such as IL-1β, tumor necrosis factor (TNF) α, and IL-6 in the context of LPS stimulation than control cells (Figures 4E and 4F). Similar results were obtained in bone-marrow-derived macrophages (BMDMs).
transfected with siMOZ (Figure S1B). However, the anti-inflammatory factor IL-10, representing the alternative activation (M2) phenotype, was higher in shMOZ cells (Figures 4E and 4F). An oxidative burst, producing various highly reactive oxygen derivatives that are critical components of the antimicrobial repertoire of macrophages, can be induced by many external stimuli, such as bacteria, fungi, and viruses (Forman and Torres, 2002). To investigate whether MOZ regulates antimicrobial reactions in macrophages, we first examined the LPS-induced oxidation level using the classical agent dichloro-dihydro-fluorescein diacetate (DCFH-DA) for indicating the amount of ROS. Consistent with reduced iNOS expression in shMOZ cells (Figure S1C), MOZ knockdown significantly suppressed the production of ROS in macrophages (Figure 4G).

To assess the role of MOZ in macrophage phagocytosis, we constructed GFP-expressing bacteria using a prokaryotic expression system that was added to the culture media of macrophages. As shown in Figures 4H and S1C, lower GFP intensity was observed in shMOZ cells, suggesting that the loss of MOZ attenuated the ability of macrophages to undergo phagocytosis. The bactericidal reaction, as a sequence of events occurring in M1 macrophages, is critical for protecting the body from infection (Banerjee et al., 2013). Here, we found that MOZ knockdown significantly diminished the bactericidal activity of macrophages, consistent with the effects of MOZ on M1 macrophage functional modulation. Taken together, these data demonstrate that MOZ regulates the cellular functions associated with the M1 phenotype.

**MOZ Positively Regulates the TLR Signaling Pathway and Histone Modification Events**

Macrophage priming with LPS initiates multiple signal transduction events, including NF-κB, mitogen-activated protein kinase (MAPK), and PI3K/AKT, which are ultimately converted into cytokine production and M1 polarization (Wang et al., 2014). To investigate whether MOZ affects the signaling events downstream of TLR4/LPS, we assessed the phosphorylation level of AKT, p65, Erk, p38, and JNK in LPS-stimulated macrophages. As shown in Figure 5A, knockdown of MOZ slightly affected the phosphorylation level of p65, but significantly decreased the phosphorylation of AKT and MAPKs. However, MOZ did not appear to regulate the expression of total protein of Erk, p38, JNK, and AKT, as evidenced by the similar bands observed in shMOZ and control cells (Figure 5A). In addition, reporter assays provided the same results with regard to the attenuation of AKT and MAPK signals in shMOZ cells, whereas MOZ did not affect the activity of NF-κB or STAT3 (Figure 5B). Thus these results suggest that MOZ regulates AKT and MAPK activity, which are crucial for M1 polarization.

Previous research has shown that MOZ can acetylate lysine-9 residues in histone H3, resulting in condensed chromatin transforming into a more relaxed structure that is associated with greater levels of gene transcription (Snyder and Glass, 2000). To determine whether MOZ regulates the histone acetylation involved in M1 polarization, we examined the acetylation level of H3 histone in LPS-stimulated shMOZ or control cells. As shown in Figure 5C, the level of acetylated and phosphorylated H3 histone was clearly reduced in shMOZ cells. In addition, chromatin immunoprecipitation assays suggested that acetylated histones binding to the enhancer of pro-inflammatory genes including IL1B and TNFA were significantly reduced in shMOZ cells (Figure 5D). In line with this finding, decreased IL-1β production was found in shMOZ cells incubated with LPS for different periods of time (Figure 5E). Thus these results demonstrate that MOZ positively regulates multiple pathways and H3 histone acetylation to promote M1 polarization.

**MOZ Is a New and Reliable Target of miR-223**

The literature indicates that miR-223 is highly expressed in myeloid cell lineages and participates in AML progression (Kuchenbauer et al., 2010). To investigate the role of miR-223 in AML, we first analyze the expression of miR-223 in AML cases based on TCGA database. Contrary to MOZ, miR-223 expression was expressed in M4/M5 patients or >20% group (Figures S3A and S3B), and low expression of miR-223 was associated with favorable prognosis (Figure S3C). To detect the expression of miR-223 or its precursor in patients with clinical AML, we designed specific primers to amplify a fragment containing miR-223, as described in Figure S5A. In accordance with previous reports, we found that the precursor or mature miR-223 was more highly expressed in AML samples...
or THP-1 cells than in healthy samples (Figures 6A and 6B). Besides, integrated data from TCGA suggested that miR-223 had inverse correlation with MOZ expression (Figure S3D). To confirm this, we examined miR-223 and MOZ expression during monocyte development. As shown in Figure 6C, miR-223 was decreased, whereas MOZ was up-regulated during PMA-, granulocyte-macrophage colony-stimulating factor (GM-CSF)-, or macrophage colony-stimulating factor (M-CSF)-induced monocyte-to-macrophage differentiation. Furthermore, we established miR-223 overexpression (designated as pre-miR-223) or knockdown (designated as sponge) cell lines (Figure S3E). Overexpression of miR-223 decreased the nuclear-cytoplasmic ratio, whereas knockdown of miR-223 increased the ratio in U937 (Figure S3J). Meanwhile, overexpression of miR-223 resulted in increased growth of U937 in soft agar containing 8 μg/mL cisplatin, whereas less growth colonies were observed after miR-223 knockdown (Figure S3K). Similar results were obtained from the CCK8 assay, which was used for analyzing the cell viability of parental U937 and the transfectants cells treated with chemotherapeutic drugs (Figures S3H and S3I). Also, overexpression of miR-223 in cells led to lower proliferation activity, whereas miR-223 knockdown promoted U937 proliferation (Figure S3L). All together, these results suggest that miR-223 promotes AML progression and inhibits monocyte differentiation.

In macrophages, miR-223 has been shown to suppress the expression of M1 phenotype and promote transition to the M2 phenotype (Li et al., 2010; Haneklaus et al., 2012). In agreement with previous reports, we found that miR-223 was also markedly reduced during LPS-induced M1 polarization, whereas M2

![Figure 4. Loss of MOZ Diminishes M1 Activation of Macrophages](image-url)
polarization induced elevated miR-223 (Figure 6D). Considering the expression pattern of MOZ and miR-223 in AML or stimulated macrophage, we speculated that MOZ may be a potential target of miR-223. To this end, we analyzed the 3'UTR of MOZ sequence using TargetScan 5.1, and we found that miR-223 was one of the conserved miRNAs targeting MOZ mRNA (Figure 6E). To validate this finding, we transfected macrophages with either synthesized miR-223 mimics or miR-223 inhibitor mimics and their corresponding scrambled controls for determining the effect on MOZ expression. As shown in Figures 6F and 6G, we found that miR-223 inhibited MOZ expression, whereas antagonizing miR-223 function by miR-223 inhibitor mimics enhanced MOZ expression. Overexpression of the precursor transcript of miR-223 resulted in reduced MOZ (Figures S3F and S5E). Besides, MOZ expression was up-regulated in U937 transfected with miR-223 sponge (Figure S3G). Reporter gene assays also demonstrated that MOZ was a validated target of miR-223 (Figures 6H and 6I). To assess whether miR-223 shares functional properties with MOZ in macrophages, BMDM cells were transfected with miR-223 mimics, siMOZ, or scramble RNA and were then examined for bactericidal activity. As shown in Figure 6J, like siMOZ, miR-223 mimics significantly suppressed bactericidal activity, suggesting that MOZ mediates the effect of miR-223 on M1 polarization.

MOZ Negatively Regulates miR-223 Expression

The critical role of PU.1 in regulating miR-223 expression has been well documented (Fukao et al., 2007). MOZ has also been reported to be a co-activator of PU.1 or RUNX1, participating in the regulation of some genes (Katsumoto et al., 2006; Kitabayashi et al., 2001). To examine whether MOZ contributes to the regulation of miR-223 in macrophages, we compared the transcriptional level of precursor or mature miR-223 in shMOZ and control cells. As expected, knockdown of MOZ enhanced the expression of miR-223 (Figures 7A and 7B and S5B/C/D). To uncover the mechanism underlying the regulation of miR-223 by MOZ, the 5'-flanking genomic region of the proximal promoter of pri-miR-223 was analyzed using JASPAR software (version: JASPAR, 2014 serve). We found a putative RUNX1-binding motif close to the confirmed PU.1-binding element on the conserved cis-acting regulatory element of the pri-miR-223 gene, and a set of reporter vectors were generated as shown in Figure 7C. Relative reporter activity for each reporter vector was examined using a dual luciferase reporter assay in shMOZ or control cells. As shown in Figure 7D, pri-miR-223–76 reporter activity was significantly higher in shMOZ cells, and a mutated putative RUNX1-binding site still exhibited enhanced activity in shMOZ cells. However, when mutation was
Figure 6. miR-223 Down-regulates MOZ Expression

(A) Total RNA was extracted from isolated and purified monocytes from AML patients or healthy volunteers following to be reverse transcribed into cDNA by using random primers or stem-loop RT primers. The cDNA was used for determining the expression precursor or mature miR-223 by using qPCR. (B) THP-1 cells or monocytes from healthy volunteers were lysed by Trizol reagent, and total RNA was reverse transcribed into cDNA. Precursor miR-223 was examined by using qPCR.

(C) miR-223 and MOZ expressions were monitored during PMA-, GM-CSF-, or M-CSF-induced monocyte differentiation at the indicated time points by qPCR.

(D) Induced human macrophage derived from monocytes (MDMs) were treated with LPS (100 ng/mL) or IL-4 (20 ng/mL) for the indicated time periods, and miR-223 and MOZ expressions were then determined by qPCR.

(E) Schematic representation of the MOZ gene and the conserved regulatory elements in the 3' UTR.

(F and G) RAW264.7 cells were transiently transfected with 20 nM miR-223 mimics or control mimics and 20 nM miR-223 inhibitors or control inhibitors. At day 3 after transfection, the cells were collected and analyzed for MOZ expression by qPCR (F) or western blot (G).

(H and I) The alignment of miR-223 and its target sites in the 3' UTR of the human MOZ gene and the introduced mutated nucleotides for the reporter assay are shown (H). A total of 200 ng psiCheck2-hMOZ-3' UTR-WT or psiCheck2-hMOZ-3' UTR-mut (hMOZ-3' UTR or hMOZ-3' UTR-mut) was co-transfected with 200 ng miR-223 overexpression plasmids into HEK293 cells. After 24 hr, Renilla luciferase activity was measured and normalized to firefly luciferase activity (I).

(J) RAW264.7 cells were transfected with 20 nM siMOZ, miR-223 mimics, or scramble small interfering RNA. At day 3 after transfection, live E. coli was added to the media for another 4 hr, and the supernatant was then diluted and cultured on Luria broth agar plates at 37°C for 16 hr. The bacterial colonies were enumerated, and the colony-forming unit of E. coli from supernatants was determined.

Data are pooled from at least three independent experiments. Data show the mean ± SEM. *p < 0.05, **p < 0.01, ***p < 0.001; NS, not significant.
introduced into the PU.1-binding sites, the pri-miR-223PU.1/C0 del reporter activity was considerably reduced and there was no difference between shMOZ and control cells. Furthermore, to exclude the possibility that MOZ regulates PU.1 expression, which sequentially affects miR-223, we examined the expression of PU.1 and RUNX1 at the transcript level in shMOZ cells. As shown in Figure 7E, there was no significant alteration of PU.1 or RUNX1 in shMOZ cells. Collectively, these results demonstrate that MOZ suppresses miR-223 expression by regulating the promoter activity of pri-miR-223, and that the PU.1-binding element on the promoter is critical for this regulation. The details of the underlying mechanism remain to be elucidated in future studies.

DISCUSSION

Previous studies have shown that MOZ fusion proteins, including MOZ-TIF2, MOZ-CBP, and MOZ-NCOA3, are associated with myeloid cancer progression (Deguchi et al., 2003; Giles et al., 1998; Esterjties et al., 2008). However, the precise role of MOZ in AML pathogenesis is not well documented. Mutant analysis of the COSMIC database showed that the somatic mutation rate of MOZ is low (0.37% in hematopoietic and lymphoid tissues). Therefore it is unlikely that somatic mutation is attributable to the deregulation of MOZ in AML progression, although we cannot exclude the deregulation of MOZ fusions in partial

Figure 7. MOZ Functions as a Repressor of the miR-223 Gene

(A and B) MOZ knockdown up-regulated miR-223 expression. Precursor or mature miR-223 level was analyzed in RAW264.7 shMOZ or control cells (A) and THP-1 shMOZ or control cells (B) by qPCR.

(C) Schematic representation of different miR-223 promoter constructs used for reporter assays.

(D) Indicated reporter plasmids were co-transfected with pRL into shMOZ or control cells. After 24 hr, the firefly luciferase activity was measured and normalized to the Renilla luciferase activity.

(E) PU.1 and RUNX1 levels were determined in shMOZ or shcon cells using qPCR. Data are presented as the mean ± SEM from three independent experiments. Similar results were obtained in three independent experiments. *p < 0.05, **p < 0.01, ***p < 0.001; NS, not significant.
patients. In addition, integrated analysis of AML in TCGA or GEO cohorts indicated that MOZ expression is the lowest in the M5 subgroup of AML and is associated with poor prognosis. Therefore combined with the observation that MOZ is up-regulated during normal monocyte development, the aberrant expression of MOZ in AML indicates that MOZ expression is strongly related to monocyte fate. A bioinformatic approach to the analysis of the promoter of the MOZ gene predicated a list of TFs with high mutation rates in AML pathogenesis, providing valuable insight into how MOZ is deregulated in AML. In our present work, we found that MOZ knockdown blocked PMA-induced monocyte differentiation and conferred apoptotic resistance to chemotherapeutic-agent-induced cell death. Thus our results reveal that loss of MOZ facilitates the neoplastic transformation of monocytes.

Derived from monocytes, macrophages undergo different activation pathways, categorized as the M1 and M2 phenotypes, in response to different external stimuli. Bacteria- or LPS-induced M1 polarization increases iNOS expression and promotes inflammatory responses mediated by up-regulated pro-inflammatory factors, whereas the M2-polarized phenotype induced by IL-4 enforces Arg-1 and IL-10 expression associated with anti-inflammatory reactions (Italiani and Boraschi, 2014). In this study, polarized M1 macrophages showed increased MOZ expression, whereas MOZ was down-regulated in IL-4-induced M2 macrophages. Knockdown of MOZ markedly attenuated the M1 activation of macrophages, as demonstrated by the reduced pro-inflammatory expression and impaired capability of phagocytosis and bacterial killing in MOZ-silenced macrophages. Together, these data suggest that MOZ plays a role in controlling the plasticity of macrophage polarization. Therefore any deregulation of MOZ could diminish the ability of macrophages to adjust their differentiation state, which is necessary for macrophages to mount a proper response to environmental cues. Conceivably, a failure of proper polarization will lead to pathologic conditions.

Intrinsic multi-faceted pathways, including the PI3K/AKT, MAPK, and NF-κB pathways, play important regulatory roles in macrophage polarization (Luyendyk et al., 2008; Kang et al., 2008; Liu et al., 2014). Upon stimulation, the receptors convert external signals into an intracellular cascade of reactions that ultimately lead to downstream transcriptional events. AKT1 kinase has been reported to regulate mitochondrial ROS production in macrophages, which induces mitophagy and modulates apoptotic resistance (Larson-Casey...
et al., 2016). In our experiments, we found that the loss of MOZ reduced phospho-AKT in response to LPS stimulation, which could explain the lower level of ROS in MOZ knockdown cells, at least partially. MOZ has been demonstrated as a co-activator for NF-κB in regulating NF-κB-dependent promoters, and the physical interaction between p65 and MOZ has been proved in vitro (Chan et al., 2007). However, in macrophages, we did not find that MOZ affected the activity of NF-κB in the context of LPS stimulation. The discrepancy between this previous study and ours could be explained by the different cell systems used (macrophage versus tumor cell lines), and the relatively low affinity of MOZ for p65 exhibited in earlier studies does not definitively show that this functional complex exists, especially in immune cells. MAPKs have at least three distinct signaling modules, including Erk, JNK, and p38 kinases that can phosphorylate the TFs c-Jun or c-Fos and induce the activation of c-Jun/c-Fos (AP-1) to shape M1 macrophage polarization (Liu et al., 2014). In our work, the impaired activity of Erk and JNK kinases, as well as the attenuated AP-1 activity, in MOZ knockdown cells suggested that MOZ could regulate the MAPK pathway to control the plasticity of macrophage polarization. However, the underlying mechanism of MOZ-mediated AKT or MAPK pathway regulation remains to be studied in the future.

Epigenetic modification plays a pivotal role in controlling the plasticity of macrophage polarization. The effector molecules involved in epigenetic regulation, including CBP, MOF, HDAC3, and JMJ2D, have been proved to be important in regulating macrophage differentiation states (Ivashkiv, 2013). MOZ has been reported to acetylate histones H3 and H4, resulting in the chromosome configuration transformation that is required for gene transcription. Indeed, we also found that MOZ regulated the phosphorylation and acetylation of H3, which is critical for pro-inflammatory factor expression. Although detailed information about how MOZ regulates pro-inflammatory factor expression (i.e., whether MOZ-mediated signal transduction and epigenetic modification are synergistic or independent for this process) remains to be explored, our data clearly demonstrate that MOZ plays multi-functional roles in macrophage activation.

miR-223, which is specifically expressed in the myeloid compartment, plays a pivotal role in limiting inflammation to prevent collateral damage during infection and in preventing oncogenic myeloid transformation. Many myeloid-related master regulators have been identified as the targets of miR-223, including C/EBPα, E2F1, IGF1R, IKKα, NFI-A, and STAT3. Some of these targets are responsible for granulopoiesis. For example, E2F1 and NFI-A have been reported to be suppressed by miR-223 at the posttranscriptional level, as well as act as repressor of miR-223 gene during granulocyte differentiation (Fazi et al., 2005; Pulikkan et al., 2010). Also, IKKα and C/EBPα as the targets of miR-223 have been identified to be critical for the differentiation of macrophage or DC (Zhou et al., 2015). Zhou et al. have reported that the miR-223/C/EBPα axis is essential for intestine homeostasis mediated by macrophage or DC. In this article, the authors have demonstrated that miR-223 deletion significantly promoted the differentiation and activation of monocyte-derived dendritic cell (MoDC). Of note, decreased Ly6C⁺ monocytes, which differentiate into intestinal CX3CR1high macrophage, and the tendency of increased granulocyte in miR-223-deficient mice suggests that the miR-223/C/EBPα axis has an important role in deciding the direction of differentiation from bone marrow (BM) granulocyte and monocyte precursors (GMPs) as reported by Zhou et al. Also, in this article, the authors find that the number of bone-marrow-derived inflammatory DCs from miR-223−/− mice induced by GM-CSF and IL-4 in vitro is much higher than from wild-type mice, suggesting that miR-223 suppresses monocyte-to-DC development. However, whether miR-223 affects monotype-to-macrophage differentiation is not mentioned by Zhou et al. Another report has shown that miR-223 is decreased during macrophage differentiation, releasing repression of its target IKKα (Li et al., 2010). In addition, miR-223 has been proved to diminish M1 macrophage polarization by targeting STAT3 (Chen et al., 2012). In line with these reports, our results showed that miR-223 was decreased during monocyte differentiation and M1 polarization. Also, like E2F1, MOZ, as a target of miR-223, was found to be a repressor of miR-223 expression. Our initial work provides a new auto-regulatory loop mediated by MOZ and miR-223 in monocyte development and macrophage activation. As mentioned above, these identified targets of miR-223 have their distinct roles in myeloid cell functions. Although we cannot rule out the possibility of other targets of miR-223 that are involved in miR-223-mediated monocyte differentiation and macrophage activation in our study, the unique role of MOZ in monocyte development provides a reliable evidence for the dysregulated miR223/MOZ blocking the normal development and promoting AML (M4/M5) pathogenesis, at least partially. And whether these targets have cross talking in miR-223-derived differentiation or inflammation remain to be uncovered in future works (Figure 8).
A previous report has shown that MOZ regulates miR-223 expression in the context of sulfatide stimulation in hepatocellular carcinoma cells (Dong et al., 2014). In this article, sulfatide has been shown to induce the disassociation of MOZ from the promoter of pre-miR-223 gene and down-regulation of miR-223 expression. However, there is no direct evidence about MOZ attachment on the promoter being essential for miR-223 gene transcription. Whether MOZ regulates miR-223 directly in hepatocellular carcinoma (HCC) by affecting the acetylation of histone or C/EBPα needs to be uncovered in the future. In our study, MOZ suppresses miR-223 expression through conserved PU.1-binding sites in the promoter of miR-223, mechanistically. Although MOZ has been demonstrated to bind with PU.1 to facilitate PU.1-dependent transcription activity in vitro (Katsumoto et al., 2006), the lack of further investigation of this process in vivo or in other cell systems cannot exclude alternative methods by which MOZ regulates PU.1-mediated transcriptional events. Based on the finding that PU.1 expression is not affected by MOZ, we consider two possible mechanisms for MOZ in regulating miR-223: one is through the inhibitory effect of MOZ on PU.1-mediated gene transcription through physical interaction and the second is by affecting other modulators involved in PU.1-initiated transcription assembly on the miR-223 promoter that are affected by MOZ. Of course, the detailed underlying mechanism for how MOZ regulates miR-223 remains to be elaborated in subsequent work.

Limitations of the Study
We want to knock out MOZ in U937 or THP-1 using CRISPR-Cas9 system; however, the efficiency of transfection is very low and we do not obtain the positive clones after several rounds of screening. Also, we try to establish AML mouse model using NOD-SCID mice to make our conclusions more convincing. However, the model in which mice were injected with THP-1 or U937 according to the well-documented protocol as reported in other literatures failed. Besides, although we had proved connection between MOZ and miR-223, we could not distinguish which is the initiator (did MOZ down-regulation in AML result in increased miR-223 or did miR-223 up-regulation result in decreased MOZ) in the regulation loop in AML pathogenesis. Another limitation is that we cannot prove how MOZ regulated signaling pathways in macrophage. We do not know whether MOZ-regulated signaling pathways resulted in cytokine expression, or MOZ regulated the cytokine expression, which induces the activation of signaling pathways.

METHODS
All methods can be found in the accompanying Transparent Methods supplemental file.

SUPPLEMENTAL INFORMATION
Supplemental Information includes Transparent Methods, five figures, and one table and can be found with this article online at https://doi.org/10.1016/j.isci.2018.12.016.

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AUTHOR CONTRIBUTIONS
Conceptualization, M.J. and J.Z.; Methodology, M.J. and J.Z.; Investigation, M.J., J.Z., L.Q., and Y.M.; Writing – Original Draft, M.J. and J.Z.; Writing – Review & Editing, M.J.; Funding Acquisition, M.J. and J.Z.; Resources, H.L. and R.L.; Supervision, M.J. and L.Q.

DECLARATION OF INTERESTS
The authors declare no competing interests.

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Supplemental Information

MOZ Forms an Autoregulatory Feedback Loop with miR-223 in AML and Monocyte/Macrophage Development

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Figure S1 Knockdown of MOZ blocks monocyte differentiation, Related to Figure 2.

(A) THP-1 was treated with 1,25(OH)\textsubscript{2}-VitD\textsubscript{3} (VD3) for indicated periods and then the expression of MOZ was determined by Q-PCR. (B) U937 cells were cultured with PMA or VD3 for indicated times and MOZ expression was examined by Q-PCR. (C) THP-1 was treated with PMA for 24 hours and subsequently be analyzed for detecting the expression of CD11b and CD14 using flow cytometry. (D) The efficiency of MOZ-knockdown in U937 was determined by Q-PCR. (E-I) Knocking down MOZ in U937 resulted in blocking differentiation. Cells numbers were counted at indicated time points for calculating the proliferate rate using trypan blue staining to exclude the dead cells (E). shMOZ and shcon cells were treated with or without cisplatin (30 \( \mu \text{g/ml} \)) for 24 hours and then flow cytometry was used for analyzing cells apoptosis which were stained with PI/Annexin V (F). Soft agar assay was used to determine the growth of shcon or shMOZ cells treated with cisplatin for 15 days. The clones were photographed by microscope (G). The morphologic analysis of shcon and shMOZ cells was used with Wright-Giemsa staining (H). The morphology of shcon or shMOZ treated with or without PMA for 24 hours was analyzed by light microscope (left and middle), PMA-stimulated cells were treated with DCFH-DA (10 \( \mu \text{M} \)) for 4 hours and subsequently to be analyzed by fluorescence microscope (I, right). Data represent the mean ±SEM from three independent experiments. Similar results were obtained in three independent experiments. *\( P<0.05 \); **\( P<0.01 \); ***\( P<0.001 \), NS, not significant.
Figure S2 MOZ inhibitor increases the activity of apoptosis resistance to cisplatin induced death, Related to Figure 2.

(A) U937 was treated with MOZ-IN-3 (3μM) for indicated periods and then the miR-223 or MOZ expression were determined by Q-PCR. (B) U937 which was pre-treated with or without MOZ-IN-3 for 15 days was stimulated with PMA (10 μg/ml) or 1,25OH-VitD3 (100 nM) for 24 hours or 48 hours, respectively. Wright-Giemsa staining were used to analyze the morphology of U937. (C-D) U937 treated with or without MOZ-IN-3 was added with cisplatin(C), cytarabine and paclitaxel (D) to analyze the apoptosis rate (C) or cell viability by using flow cytometry or CCK8 assay, respectively. Data represent the mean ±SEM from three independent experiments. Similar results were obtained in three independent experiments. *P<0.05; **P<0.01; ***P<0.001, NS, not significant.
Figure S3 High expression of miR-223 is associated with poor prognosis, Related to Figure 1 and Figure 6.

(A) Integrate analysis of miR-223 expression in different subtypes of AML based on TCGA database. (B) Integrate analysis of miR-223 expression in AML cases which were subdivided into 3 groups based on the proportion of monocytes (=0, <20% and >20%). (C) Integrate analysis of miR-223 expression in AML subgroups with poor, intermediate or advanced prognosis. (D) Correlation analysis between MOZ and miR-223 expression. (E) Schematic representation of design for miR-223 overexpression or knockdown. (F-G) Established miR-223 overexpression or knockdown cells were analyzed for the MOZ expression by using Q-PCR. (H-I) miR-223 overexpression or knockdown cells and their corresponding control cells were treated with chemotherapeutic drugs at indicated concentration for 24 hours and then analyzed using CCK8 assay. (J) Wright-Giemsa staining analyzed the morphology of miR-223 overexpression or knockdown cells. (K) miR-223 overexpression or knockdown cells and their corresponding control cells were treated with cisplatin (8 μg/ml) and then seeded into soft agar for 15 days, the visible colonies were photographed by light microscope. (L) miR-223 overexpression or knockdown cells and their corresponding control cells were counted using trypan blue staining and the number of live cells were calculated for growth curve. Data are presented as the mean ±SEM from three independent experiments. Similar results were obtained in three independent experiments. *P<0.05; **P<0.01; ***P<0.001, NS, not significant.
Figure S4 Knockdown of MOZ dampens the M1 activity of BMDM, Related to Figure 4.

(A) BMDMs were transfected with 20 nM siMOZ or control siRNA. At day 3 after transfection, the efficiency of knockdown was determined. (B-C) BMDMs transfected with siMOZ or scramble siRNA were stimulated with LPS (100 ng/ml) or IL-4 (20 ng/ml) for the indicated time periods and subjected to analysis of the levels of M1 markers (B) or the M2 marker Arginase 1 (C) by Q-PCR. (D) siMOZ or control BMDMs were pretreated with LPS for 12 hours. After stimulation, live E. coli was added to the media for another 2 hours, and the CFUs of the supernatant were then determined. (E) siMOZ or control BMDMs were treated with TLR agonists at the indicated concentrations together with GFP-bacteria for 4 hours and subjected to FACS analysis for GFP intensity. Data represent the mean ±SEM from three independent experiments. Similar results were obtained in three independent experiments. *P<0.05; **P<0.01; ***P<0.001, NS, not significant.
Figure S5 MOZ acts as a target of miR-223 and a repressor for the miR-223 gene, Related to Figure 7.

(A) Schematic representation of the miR-223 gene. (B) RAW264.7 cells were transiently transfected with overexpression plasmids of miR-223 (pcDNA3.1-pre-miR-233) or control plasmids. After 48 hours, MOZ expression was measured by Q-PCR. (C) 20 nM siMOZ or scramble siRNA was transfected into BMDCMs. After 48 hours, the precursor of miR-223 was detected by Q-PCR. (D) Stably transfected THP-1 cells with either shPU.1 or control shRNA were examined for miR-223 expression levels by Q-PCR. Data represents the mean ±SEM from three independent experiments. Similar results were obtained in three independent experiments. *P<0.05; **P<0.01; ***P<0.001, NS, not significant.
**Table S1** The primers used in our present works are shown, Related to Figure 2, Figure 4, Figure 6 and Figure 7.
**Transparent Methods**

**Mice and reagents**

C57BL/6 mice were purchased from SIPPR-BK Experimental Animal Ltd. Co. (Shanghai, China). Experiments and animal care were performed according to protocols approved by the University of Science and Technology of China (USTC) Institutional Animal Care and Use Committee. Lipopolysaccharide (LPS), polyI:C and PAM3CK4 were purchased from Sigma (St. Louis, MO). Recombinant human and mouse GM-CSF and M-CSF were from PeproTech (Rocky Hill, NJ). 1,25OH-VitD3 was purchased from Sigma (St. Louis, MO). MOZ-IN-3 was from MCE (MedChem Express (Monmouth Junction, NJ, USA). MiR-223 mimics and control, miR-223 inhibitors and inhibitor control, MOZ small interfering RNA (siRNA) and scrambled control siRNA were purchased from GenePharma (Shanghai, China). CCK8 kit and Wright-Giemsa solution was obtained from Solarbio (Beijing, China). Primary antibodies against p65/phospho-p65 (S560), AKT/phospho-AKT (S473), p38/phospho-p38, Erk/phospho-Erk1/2, and JNK/phospho-JNK were obtained from Cell Signal Technology (Danvers, MA). Antibodies against MOZ were purchased from Abcam. Antibodies against GAPDH and Ac-lysine were purchased from Santa Cruz Biotechnology (Santa Cruz, CA). Phosphorylated and acetylated Histone H3 antibody was purchased from Genetex Inc. (California, USA). HRP-conjugated secondary antibodies were obtained from Biolegend. FITC-coupled and PE-coupled CD34 or CD38 antibodies, respectively, for flow cytometry were obtained from BD Biosciences. ELISA kits detecting IL-6 and IL-1β were purchased from Lianke Ct.

**Monocyte isolation and cell culture**

The study was approved by the institutional ethics board of the University of Science and Technology of China and conducted in accordance with the Declaration of Helsinki. Human monocytes were isolated from the peripheral blood of healthy adult volunteers using consecutive Ficoll-Hypaque and Percoll density gradient centrifugation as reported previously (Todd and Schlossman, 1982). The purity of monocytes was ~94% (± 2%) based on flow cytometry analysis of CD14. Human monocyte-derived macrophages were generated by culturing monocytes in the presence of GM-CSF as described previously (Cassol et al., 2009). Briefly, monocytes were incubated at 5×10^5 cells/ml in RPMI 1640 medium (RPMI 1640 plus 2 mM glutamine, 25 mM HEPES, 100 U/ml penicillin, and 100 mg/ml streptomycin) containing 10% FBS and 100 ng/ml recombinant human or mouse GM-CSF at 37°C in a 5% CO_2 incubator for 5 days. BMDMs were derived from the bone marrow of C57BL/6 mice. In brief, after lysis of red blood cells, bone marrow cells were cultured in DMEM media containing 10% FBS and 20 ng/ml M-CSF for 5 days to establish macrophages. RAW264.7 and HEK293 cells were maintained in DMEM supplemented with 10% FBS, 100 U/ml penicillin, and 100 mg/ml streptomycin. THP-1 cells were cultured in RPMI 1640 supplemented with 10% FBS and standard concentrations of penicillin-streptomycin. To induce M1 polarization, macrophages were cultured in the same cytokine mixture in the presence of LPS (100 ng/ml) for an indicated period of time at 37°C in a 5% CO_2 incubator.

**Plasmids and transfection**

The library of shRNA used for producing viral-packaged interfering RNA for target genes was obtained from Sigma. The protocol of lentiviral particle production was performed as previously
described (Hornung et al., 2009). For generating stable cell lines, lentiviral particles were added together with Polybrene to 6-well plates that had been seeded with \(8 \times 10^5\) cells, and the final concentration of Polybrene was 8 μg/ml. Cells were incubated overnight, and the lentivirus/Polybrene mixture was replaced with fresh puromycin-containing culture media every 3-4 days until resistant colonies could be identified. For knocking down MOZ with synthetic siRNA in BMDMs, \(6 \times 10^6\) cells were seeded into 6-well plates. Lipofectamine 2000 was used to deliver siRNA into cells as per the manufacturer’s recommendations.

**Luciferase assay**

A 1-kb fragment containing the predicted miR-223 binding element in the MOZ 3'UTR was obtained by PCR amplification using human genomic DNA as a template and was then cloned into the Bgl-II and EcoRI sites of the psiCheck2-Reporter Luciferase vector. The resulting construct was designated hMOZ-3'UTR. For mutant MOZ 3'UTR construction, nucleotides in the 3'UTR of the MOZ gene that matched the miR-223 “seed sequence” were mutated by site-directed mutagenesis using site-directed mutagenesis kits (TransGen, Biotech). The resulting construct was designated hMOZ-3'UTR-mut. The sequences of the primers used for PCR amplification are listed in Table S1. hMOZ-3'UTR or -mut was cotransfected with miR-223 overexpression plasmids using Lipofectamine 2000 reagent (Invitrogen). At 24 hours after transfection, the luciferase activity was determined using the Dual Luciferase Assay System (Promega). For analysis of the promoter activity of miR-223, a -761 to +21 bp fragment (relative to the translational start site) containing the confirmed PU.1 binding sites and predicted RUNX1 binding site was amplified from mouse genomic DNA using high-fidelity PCR and cloned into pGL3-basic vectors. pri-miR-223\(^{PU.1\text{-del}}\) and pri-miR-223\(^{RUNX1\text{-mut}}\) reporter constructs were identical to the wild-type construct except that the PU.1-binding sites were deleted or the predicted RUNX1-binding site was inactivated by substitution of CAGAA at positions +59 to +54, respectively. The related primers are in Table S1. Established reporter constructs were transiently co-transfected with pRL plasmids into RAW264.7 cells, and the lysed supernatant was measured using a dual-reporter Luciferase assay (Promega) to determine the luciferase activity.

**Bacterial killing assay**

A bacterial killing assay was performed as described previously (Banerjee et al., 2013). Briefly, 0.1 \(\times 10^6\) CFU/ml of *E. coli* (BL21DE3pLysS) was added to media in 48-well plates and incubated at 37°C for 4 hours. The supernatants were then serially diluted 10 times, and 20 μl of each diluted supernatant was seeded on LB-agar plates. The plates were incubated overnight at 37°C, and bacterial colonies were counted after 16 hours.

**Bacterial phagocytosis assay**

The bacteria for this assay were previously transformed with pET28-GFP vectors. GFP-expressing bacterial clones were co-cultured with macrophages in 48-well plates. After incubation for 4 hours at 37°C in a 5% CO\(_2\) incubator, the macrophages were washed twice with cold PBS, and a flow cytometry assay was then performed.

**Soft agar and CCK8 assay**

Colony formation assay in soft agar were done according with standard protocols. In brief, pre-
warmed 2×RPMI-1640 containing 20% FBS, 200 U/ml penicillin, 200 µg/ml streptomycin and equivalent melted 1.2% SeaPlaque Agarose (Lonza) solution were mixed and transferred into a well (3 ml per well) in a 6-well plates, which was then incubated at room temperature for 30 min to allow the bottom agar layer to solidify. Cell suspension (5×10^3 per well) in RPMI-1640/10% FBS was mixed with equivalent 2×RPIM-1640 containing 20% FBS and same volume of 1.2% agar. The mixture was then plated on the bottom agar layer and the top agar was solidified at room temperature before the plates being put into CO2-incubator. After about 15 days, the visible colonies were counted by light microscope.

CCK8 assay was performed with standard protocol. Briefly, parental cells (U937) and their transfectants were plated in 96-well plates at a density of 1×10^4 per well in 200 µl of RPMI-1640 with 10% FBS. After 12 hours, different dose of chemotherapeutic drugs were added into 96-well plates as demanded for 24 hours, then 20 µl of CCK8 solution was added per well. After incubation for 2 hours at 37℃, we measured the absorbance of each well at 450 nm by using an ELISA plate reader.

Quantitative real-time PCR

Total cellular RNA was extracted using Trizol reagent (Invitrogen). A quantity of 2 µg of total RNA was used for the reverse-transcription reaction using the First Strand cDNA Synthesis kit (Invitrogen). The obtained cDNA was analyzed using real-time PCR, as previously described (Pulikkan et al., 2010). Primer sequences are listed in Table S1.

Immunoblot analysis

For preparing the whole-cell extract, cell were harvested and lysed in radioimmunoprecipitation assay buffer (50 mM Tris-HCl [pH 7.4], 1% Nonidet P-40, 0.25% sodium deoxycholate, 150 mM NaCl, 1 mM EDTA, 1 mM PMSF, 1 mg/ml aprotinin, 1 mg/ml leupeptin, 1 mM pepstatin, 1 mM Na3VO4, and 1 mM NaF). The protein concentration of lysate was quantified using a Bicinchoninic Acid Protein Assay Kit (Pierce, Rockford, IL). Cell lysates were loading onto an SDS-PAGE gel and separated by electrophoresis. After transfer of the proteins onto a nitrocellulose membrane, the target molecules were detected by Western blotting using corresponding primary antibodies, HRP-conjugated secondary antibodies, and an ECL Detection kit (Pierce).

Measurement of cytokines and chemokines

The supernatants of cultured macrophages were harvested and kept at -80℃. The concentrations of selected cytokines and chemokines were measured using an ELISA kit (Lianke) or a CBA kit (BD) following standard procedures.

Data mining and bioinformatics analysis

Related gene expression profiles in AML cases were extracted from the TCGA and GEO databases. The normalized expression value of related genes in AML cases was used for correlation analysis and survival analysis (Kaplan-Meier) using SPSS software. The analysis of transcriptional regulation of the MOZ gene was based on putative transcription factors predicated by various databases or software (PROMO 3.0, JASPAR, COMPEL, FANTOM and Ensembl) on 3 kb of sequence upstream of the proximal transcription start site of the MOZ gene. These predicated transcription factors (the setting threshold was >0.9 or 90 according to the different software used)
were clustered using DAVID 6.8, and co-expression was analyzed using cytoscape software.

Statistical analysis

All experiments were repeated three times. The data are presented as the mean ±SEM. Statistical significance was determined by Student’s t test and analysis of variance (ANOVA) using SPSS, with a P value less than 0.05 considered statistically significant.