microRNA-7 Suppresses the Invasive Potential of Breast Cancer Cells and Sensitizes Cells to DNA Damages by Targeting Histone Methyltransferase SET8*

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**Background:** How SET8 is regulated is not fully understood.

**Results:** MicroRNA-7 down-regulates SET8 and inhibits H4K20 monomethylation, suppresses metastasis of breast cancer cells, and sensitizes cells to DNA damages.

**Conclusion:** MicroRNA-7 is a negative regulator of SET8.

**Significance:** This work aids our understanding of the biological function of microRNA-7, supporting the pursuit of microRNA-7 as a potential target for breast cancer intervention.

SET8 (SET domain containing 8) is a histone H4 lysine 20 (H4K20)-specific monomethyltransferase in higher eukaryotes that exerts diverse functions in transcription regulation, DNA repair, tumor metastasis, and genome integrity. The activity of SET8 is tightly controlled during cell cycle through post-translational modifications, including ubiquitination, phosphorylation, and sumoylation. However, how the expression of SET8 is regulated is not fully understood. Here, we report that microRNA-7 is a negative regulator of SET8. We demonstrated that microRNA-7 inhibits H4K20 monomethylation and suppresses epithelial-mesenchymal transition and the invasive potential of breast cancer cells. We showed that microRNA-7 promotes spontaneous DNA damages and sensitizes cells to induced DNA damages. Our experiments provide a molecular mechanism for the regulation of SET8 and extend the biological function of microRNA-7 to DNA damage response, supporting the pursuit of microRNA-7 as a potential target for breast cancer intervention.

MicroRNAs (miRNAs) are a cluster of evolutionarily conserved, non-encoding single-stranded RNA molecules of 20–24 nucleotides. These molecules function to silence gene expression by binding to partially complementary recognition sequences of the 3’-UTRs of target mRNAs, leading to either mRNA degradation or translation inhibition (1, 2). More than 30% of human genes appear to have been under selective pressure to maintain their pairing to miRNA seeds, suggesting a role of miRNAs in posttranscriptional repression of these genes. With a growing number of oncogenes and tumor suppressor genes that have been found to be under the control of miRNAs, the importance of miRNAs in cell cycle progression, cell proliferation, genome stability, and malignant transformation is becoming increasingly recognized (3).

MicroRNA-7 (miR-7) is an intronic miRNA that resides in three different gene loci in the human genome and is conserved among all species (4). Aberrant expression of miR-7 was observed in various malignancies, including breast cancer (5, 6), glioblastomas (7), and gastric and colorectal cancers (8). Functionally, miR-7 has been characterized as a putative tumor suppressor through its suppression of the expression of multiple EGF receptor signaling-related genes, including EGF receptor as well as its downstream kinase RAF1 (9), p21-activated kinase 1, and insulin receptor substrate-2 (10). Moreover, a range of functionally important genes, including Kruppel-like factor 4 (5), Yin Yang 1 (YY1) (11), focal adhesion kinase (6), and insulin-like growth factor 1 receptor (12), have also been reported to be miR-7 targets, suggesting its profound role in malignant transformation and metastasis.

SET8 (also known as PR-Set7/9, SETD8, KMT5A), a member of the SET domain-containing methyltransferase family that specifically targets H4K20 for monomethylation, has been implicated in a diverse array of biological processes, including controlling gene transcription (13, 14), maintaining genome integrity (15, 16), regulating cell cycle progression (17, 18), and mediating DNA damage and repair (19, 20). In addition to H4K20, SET8 was also reported to methylate p53 at lysine residue 382 (p53K382me1) and to repress p53-mediated transcriptional activation of target genes (21). Previously, we reported that SET8 is functionally linked to TWIST and acts, via its H4K20 monomethylation activity, as a dual epigenetic modifier on the promoters of the TWIST target genes E-cadherin and N-cadherin (22).

Although the importance of SET8 in cell proliferation and homeostasis is well recognized and its activity is believed to be...
Targeting SET8 by microRNA-7

…tightly controlled during cell cycle through post-translational modifications, including ubiquitination, phosphorylation, and sumoylation, how the expression of SET8 is regulated is poorly understood. In this report, we identified miR-7 as a negative regulator of SET8. We showed that miR-7 inhibits H4K20 monomethylation through targeting the 3′-UTR of SET8 mRNA. We demonstrated that miR-7 suppresses epithelial-mesenchymal transition (EMT) and invasion of breast cancer cells. We found that miR-7 promotes spontaneous DNA damage and sensitizes cells to induced DNA damages.

EXPERIMENTAL PROCEDURES

Antibodies and Reagents—The source of antibodies against following proteins was as follows. Tubulin and p53 were from B & M BIOTECH Co., Ltd.; SET8, RAD51, and 53BP1 were from Cell Signaling Technology; E-cadherin, α-catenin, β-catenin, N-cadherin, and γ-catenin were from BD Biosciences; fibronectin, H4, H3K4me3, H3K9me3, H3K27me3, H4K36me3, and H4K20me1 were from Abcam; H2AX was from Millipore. Control siRNA, SET8 siRNA, and miR-7 mimics were synthesized by Shanghai GeneChem, Inc. (Shanghai, China). MiR-7 inhibitors were obtained from Dharmacon.

Cell Culture and Cell Transfection—MCF-7 and MDA-MB-231 cells were from the American Type Culture Collection (Manassas, VA). MCF-7 cells were maintained in DMEM (Hyclone) supplemented with 10% FBS. MDA-MB-231 cells were cultured in Leibovitz’s L-15 medium with 10% FBS at 37°C without CO₂. Plasmids transfection was carried out using Megatran (version 1.0, Origene) according to the manufacturer’s recommendations.

Plasmid Construction and Luciferase Assays—The wild-type and mutant SET8 3′-UTR were amplified by PCR and cloned in pmIR-REPORT (Ambion) with firefly luciferase. These PCR products were digested and ligated into the SpeI-Hind III sites of pmIR-REPORT vectors to generate a series of reporter constructs. The luciferase assays were performed in MCF-7 cells as described previously (23). MCF-7 cells treated with control, miR-7 mimics, or miR-7 inhibitors were transfected with wild-type or mutants of SET8 3′-UTR luciferase reporters together with Renilla plasmid. 48 h after transfection, the firefly and Renilla luciferases were assayed according to the manufacturer’s instructions (Promega), and the firefly luciferase activity was normalized to that of Renilla luciferase. Each experiment was repeated in triplicate. SET8 3′-UTR wt-1 (forward primer), 5′-CTTCTTCTAAAAGGAAAAATGCC-3′; SET8 3′-UTR wt-1 (reverse primer), 5′-TACAAAGCTAAAACCACAAGAAGG-3′; SET8 3′-UTR wt-2 (forward primer), 5′-GGACTAGTTCTGAAAGCAAAATGCC-3′; SET8 3′-UTR wt-2 (reverse primer), 5′-GGGCCCAGCACTTTCACTGGAGAG-3′; SET8 3′-UTR wt-3 (forward primer), 5′-GGACTAGTTCTGCAAACTAAG-3′; SET8 3′-UTR wt-3 (reverse primer), 5′-GGGCCCAGCACTTTCACTGGAGAG-3′; SET8 3′-UTR wt-4 (forward primer), 5′-ATGCAGTCAAGACTCAGCACAGCAGGAGG-3′; SET8 3′-UTR mut-2 (forward primer), 5′-TTTGTGTGCGATGACATCTTCCCTCTGCTTCTCGGA-3′; SET8 3′-UTR mut-2 (reverse primer), 5′-ATGCAGTCAAGACTCAGCACAGCAGGAGG-3′; SET8 3′-UTR mut-3 (forward primer), 5′-CCGGGCATATTTGCCATACACCCCTGGCTTTCTC-3′; SET8 3′-UTR mut-3 (reverse primer), 5′-AGAAAGCCGGAGCCTTTGCTACGTGAAATCTATGGCAGG-3′; and SET8 3′-UTR mut-4 (forward primer), 5′-ATGCAGTCAAGACTCAGCACAGCAGGAGG-3′.

RNA Isolation and Real-time Quantitative PCR—Total cellular RNAs were isolated with the TRizol (Invitrogen) and the first strand cDNA synthesis was performed with the reverse transcription system (Promega). Quantization of all gene transcripts was done by real-time quantitative PCR using Power SYBR Green PCR Master Mix and an ABIPRISM 7300 sequence detection system (Applied Biosystems, Foster City, CA) with the expression of GAPDH or U6 as the internal control. The primer pairs used were as follows: GAPDH (forward primer), 5′-CCCCACTCTCCACCTTTGCAC-3′ and GAPDH (reverse primer), 5′-CATAACAGGAATATGTGGTCG-3′; miR-7 (forward primer), 5′-CTCAACTGGTTGTGAAGTCG-3′ and miR-7 (reverse transcription primer), 5′-CCTACTGGTTGTGAAGTCG-3′; E-cadherin, 5′-CTCAACTGGTTGTGAAGTCG-3′; miR-7 (forward primer), 5′-CATAACAGGAATATGTGGTCG-3′; and miR-7 (reverse primer), 5′-CTCAACTGGTTGTGAAGTCG-3′; H2AX was from Millipore. Conjugation X-100 in PBS, blocked with 0.8% BSA, and incubated with appropriate primary antibodies followed by staining with FITC or RITC-conjugated secondary antibodies. Cells were washed four times and a final concentration of 0.1 μg/ml DAPI (Sigma) was included in the last washing to stain nuclei. Images were visualized and recorded with an Olympus FV1000S confocal microscope.

Transwell Invasion Assay—The Transwell invasion assays were performed using the Transwell chamber (Chemicon, Inc.) with a Matrigel-coated filter. MDA-MB-231 cells were transfected with control, miR-7 mimics, or miR-7 inhibitors with/without SET8 cDNA lacking 3′-UTR for 36 h. Cells were washed with PBS, fixed in 4% (w/v) paraformaldehyde, permeabilized with 0.1% (v/v) Triton X-100 in PBS, blocked with 0.8% BSA, and incubated with appropriate primary antibodies followed by staining with FITC or RITC-conjugated secondary antibodies. Cells were washed four times and a final concentration of 0.1 μg/ml DAPI (Sigma) was included in the last washing to stain nuclei. Images were visualized and recorded with an Olympus FV1000S confocal microscope.
removed using cotton swabs, and the invasive cells on the lower side were fixed, stained with 0.1% crystal violet solution, and counted using light microscope. The experiments were repeated three times.

Homologous Recombination (HR) and Nonhomologous End-joining (NHEJ) Assays—DR-GFP-U2OS and EJ5-GFP-HEK293 stable cell lines (kindly provided by Dr. Xingzhi Xu, The Capital Normal University) were transfected with control or miR-7 mimics with/without \( \text{SET8} \) cDNA lacking 3'-UTR for 6 h prior to transfection with PCBASce or pCAGGS-Ds-red (transfection indicator). After 48 h of transfection, HR or NHEJ repair efficiency was measured with the percentage of GFP-positive by FACSscan.

Clonogenic Assay—MCF-7 cells were transfected with control or miR-7 mimics with/without \( \text{SET8} \) cDNA lacking 3'-UTR. After 48 h of transfection, cells were challenged with different concentrations of VP16 or camptothecin, and then were plated in triplicate wells at 1000 cells/60-mm² dish. After 14 days, the colonies were stained with 0.1% crystal violet solution, and counted using light microscope. Colonies were defined as groups of 30 or more cells and survival potential was calculated as a percentage of colonies relative to untreated controls. The experiment was repeated three times.

RESULTS

Histone Methyltransferase \( \text{SET8} \) Is a Downstream Target of miR-7—MiR-7 is implicated in cell proliferation and apoptosis, and its expression is dysregulated in many types of tumors, including breast cancer (10). To further explore the functional role of miR-7 in pathophysiological settings, we sought to identify additional targets for miR-7. Analyses based on three public algorithms (TargetScan, PicTar, and miRanda) and the data about miR-7-downregulated genes in three cell lines (HNS, A549, and FaDu) from microarrays (9, 24) yielded several potential targets for miR-7 with high confidence scores (Fig. 1A), including \( \text{SET8} \), PSME3, POLE4, CKAP4, and CNN3, as well as RAF1, a known target of miR-7. In light of our previous observation that \( \text{SET8} \) is implicated in the development and progression of breast cancer (22), we focused our study on \( \text{SET8} \). Western blotting analysis showed a marked reduced level of \( \text{SET8} \) protein in miR-7 mimic-transfected MCF-7 cells (Fig. 1B), and further bioinformatics analysis of miRanda indicated that the \( \text{SET8} \) 3'-UTR harbors four potential miR-7 target sites (A, B, C, and D) (Fig. 1C). These sites are either completely (B and D) or partially (A and C) complementary to a heptamer motif 5'-GGAGAGC-3' that is found in the seed region of human miR-7 (Fig. 1C).

To verify the proposition that \( \text{SET8} \) is a downstream target of miR-7, reporter assays were performed in MCF-7 cells with the luciferase gene driven by either wild-type or mutated \( \text{SET8} \) 3'-UTR sequences (Fig. 1D), including the full-length and three deletions of the wild-type \( \text{SET8} \) 3'-UTR (wt-1, 2, 3, 4) as well as 4 mutants of \( \text{SET8} \) 3'-UTR (mut-1, 2, 3, 4), with each containing three point mutations (Fig. 1E). The results showed a 60% reduction in luciferase activity with wt-1 (the full-length wild-type \( \text{SET8} \) 3'-UTR) in MCF-7 cells transfected with miR-7 mimics. Consistent with our prediction, the sites B and D play a more important role in the binding of miR-7 to \( \text{SET8} \) 3'-UTR, as there were more evident decreases in luciferase activity with wt-3, wt-4 and mut-3, compared with that with wt-2 (Fig. 1F). Moreover, the binding of miR-7 to sites B and D of \( \text{SET8} \) 3'-UTR was specific, as the point mutation of corresponding sites (mut-2 and mut-4) disrupted miR-7 repressive activity on reporter gene expression, whereas mutation of sites A (mut-1) did not change luciferase activity (Fig. 1F). Consistently, treatment with miR-7 inhibitors led to a reverse trend of luciferase activity with wt-1, 3, 4, and mut-3 (Fig. 1G). Taken together, these results suggest that miR-7 binds to \( \text{SET8} \) 3'-UTR and that \( \text{SET8} \) is a downstream target of miR-7.

miR-7 Promotes \( \text{SET8} \) mRNA Degradation and Inhibits H4K20 Monomethylation—In view of the fact that miRNAs silence genes by translational inhibition and mRNA destabilization (25), we next investigated the effect of miR-7 on the expression of \( \text{SET8} \) mRNA and protein. For this purpose, MDA-MB-231 and MCF-7 cells were transfected with control or miR-7 mimics, followed by the measurement of \( \text{SET8} \) mRNA and protein expression by real-time quantitative PCR (qPCR) and Western blotting, respectively. The results showed that miR-7 overexpression led to ~40 and 60% decrease in \( \text{SET8} \) mRNA in MDA-MB-231 and MCF-7 cells, respectively (Fig. 2A), and Western blotting analysis also showed a marked reduction of \( \text{SET8} \) protein expression in cells transfected with miR-7 mimics (Fig. 2A). Consistently, treatment of the cells with miR-7 inhibitors resulted in an ~2-fold increase in \( \text{SET8} \) mRNA (Fig. 2B) and a pronounced elevation in \( \text{SET8} \) protein expression (Fig. 2B). To determine the impact of miR-7 on \( \text{SET8} \) mRNA stability, MCF-7 cells and MDA-MB-231 cells were transfected with miR-7 mimics or miR-7 inhibitors followed by treatment with transcription inhibitor actinomycin D. Real-time qPCR measurement showed that miR-7 depletion was associated with an increased \( \text{SET8} \) mRNA half-time, whereas miR-7 overexpression led to a reduced \( \text{SET8} \) mRNA half-time (from 8 to 4 h) (Fig. 2C), further supporting the notion that miR-7 represses \( \text{SET8} \) expression through destabilization of \( \text{SET8} \) mRNA.

As stated above, \( \text{SET8} \) is a member of SET domain-containing methyltransferase family that specifically catalyzes monomethylation of histone H4 Lys-20 (H4K20me1). The observation that miR-7 influences the mRNA level of \( \text{SET8} \) prompted us to investigate whether miR-7 could impact on monomethylation of H4K20. Consistent with decreased expression of \( \text{SET8} \) protein, reduction of H4K20me1 was observed in miR-7 mimic-transfected MCF-7 cells and MDA-MB-231 cells, whereas no significant difference in H3K4me3, H3K9me3, H3K27me3, H3K36me3, and total histone H4 were detected in these cells (Fig. 2D). In contrast, cells that were treated with miR-7 inhibitors exhibited increased levels of H4K20me1 (Fig. 2D), supporting the argument that miR-7 inhibits H4K20 monomethylation by promoting \( \text{SET8} \) mRNA degradation.

It was reported that \( \text{SET8} \) could also methylate p53 at lysine 382 (p53K382) and suppress p53-mediated transcription activation of target genes such as p21 protein (21). Therefore, we tested whether miR-7 could regulate the activity of p53 and the expression of its downstream genes in breast cancer cells. Western blotting analysis showed an elevation of p21 expression in miR-7 mimic-treated cells, whereas miR-7 inhibitor treatment led to a reverse trend of p21 protein but not p53 itself.
Targeting SET8 by microRNA-7

FIGURE 1. Histone methyltransferase SET8 is a downstream target of miR-7. A, the intersection of the left Venn diagram displays 25 potential targets for miR-7 predicted through PicTar, TargetScan, and miranda algorithms with high confidence scores. The intersection of the right Venn diagram displays 17 down-regulated genes (>2.0-fold) based on microarray data in miR-7-overexpressed A549, HN5, and FaDu cells. The intersection of the lower Venn diagram is an overlap between algorithm prediction and microarray data analysis, representing six candidate targets for miR-7. B, MCF-7 cells were transfected with control or miR-7 mimics for 48 h. The endogenous SET8 protein expression was measured by Western blotting. C, sequence alignment of SET8 3′-UTR between human (Homo sapiens), chimpanzee (Pan troglodytes), dog (Canis familiaris), and rat (Rattus norvegicus). The miR-7 seed sequence (5′-GGAGAC-3′) and the corresponding complementary sequences of SET8 3′-UTR are shaded. D, the diagram of luciferase reporter constructs for full-length wild-type (wt-1), truncated wild-type (wt-2, 3, 4), and mutants (mut-1, 2, 3, 4) of SET8 3′-UTR. E, sequences of wild-type and mutant target sites for miR-7 in SET8 3′-UTR are shown. miR-7 seed sequence (5′-GGAGAC-3′) is shaded. Three point mutations (underlined) predicted to abolish miRNA-mRNA binding were made in miR-7 recognition region. F, luciferase reporter assays were performed to identify the binding of miR-7 to SET8 3′-UTR. MCF-7 cells were co-transfected with negative control (NC) or miR-7 mimics (miR-7) together with the luciferase gene driven by either wild-type or mutated SET8 3′-UTR sequences, including the full-length and three deletions of the wild-type SET8 3′-UTR (wt-1, 2, 3, 4) as well as four mutants of SET8 3′-UTR (mut-1, 2, 3, 4). The normalized luciferase activity in the control group was set as relative luciferase activity. Each bar represents the mean ± S.D. for triplicate experiments. p values were determined by Student’s t test. *, p < 0.05.
G, luciferase reporter assays were performed in MCF-7 cells that co-transfected with negative control (In-NC) or miR-7 inhibitors (In-miR-7) together with the luciferase gene driven by either wild-type or mutated SET8 3′-UTR sequences, including the full-length and three deletions of the wild-type SET8 3′-UTR (wt-1, 2, 3, 4) as well as four mutants of SET8 3′-UTR (mut-1, 2, 3, 4). The normalized luciferase activity in the control group was set as relative luciferase activity. Each bar represents the mean ± S.D. for triplicate experiments. p values were determined by Student’s t test. *, p < 0.05.

(Fig. 2D). Collectively, these data indicate that miR-7 inhibits H4K20 monomethylation and promotes p53 transcriptional activity by targeting SET8.

miR-7 Suppresses EMT and Invasion of Breast Cancer Cells by Down-regulating SET8—We next investigated the biological significance of miR-7-downregulated SET8 expression. To this
in non-metastatic MCF-7 cells. Based on our previous report that SET8 promotes EMT and metastasis of breast cancer (22), it is reasonable to postulate that miR-7 also play a role in EMT and metastasis of breast cancer cells. To test this hypothesis, we examined cell morphology and expression of epithelial/mesenchymal markers in miR-7 mimic- or miR-7 inhibitor-transfected MDA-MB-231 cells by microscopy and Western blotting, respectively. Although control MDA-MB-231 cells maintained a spindle-like, fibroblastic morphology representing mesenchymal cell morphology, miR-7 mimic-transfected cells displayed a scattering and cobble stone-like appearance (Fig. 3B). Meanwhile, miR-7 inhibitor-treated cells still kept mesenchymal cell morphology and even became longer and thinner (Fig. 3B). Consistent with this, immunofluorescent microscopy showed an elevation of E-cadherin and γ-catenin staining in miR-7-overexpressing MDA-MB-231 cells compared with that in control cells, whereas vimentin, a mesenchymal marker, exhibited a reverse trend (Fig. 3C). In agreement with the results, we found that overexpression of miR-7 resulted in increases in the expression levels of epithelial markers (E-cadherin, α-catenin, β-catenin, and γ-catenin) and decreases in that of mesenchymal markers (N-cadherin, fibronectin, and vimentin), as shown by immunoblotting (Fig. 3D). Conversely, repression of miR-7 with specific inhibitors in MDA-MB-231 cells was associated with an evident derepression of the mesenchymal markers and a significant deactivation of the epithelial markers, as shown by immunoblotting (Fig. 3D). Moreover, miR-7 expressing MDA-MB-231 cells that were co-transfected with the SET8 expression construct containing SET8 coding sequence but lacking the SET8 3′-UTR, which is resistant to miR-7, attenuated the morphological alterations and epithelial/mesenchymal marker changes induced by miR-7 (Fig. 3, B–D). The restoration of SET8 expression was confirmed by immunoblotting (Fig. 3D), supporting a notion that the impact of miR-7 on EMT and expression of epithelial/mesenchymal markers in breast cancer cells is mediated by SET8.

To substantiate the role of miR-7 in tumor migration and invasion, the effect of gain-of-function or loss-of-function of miR-7 on the invasive potential in MDA-MB-231 cells was investigated by wound-healing assays. The results showed that miR-7 inhibitor-treated cells were more efficient in wound healing, whereas miR-7 mimic-treated cells were somewhat resistant to wound healing (Fig. 3E). In addition, transwell invasion assays showed that miR-7 overexpression inhibited the invasion of MDA-MB-231 cells by 4.5-fold, whereas miR-7 knockdown promoted the invasion by 3-fold (Fig. 3F). Moreover, miR-7-suppressed EMT of breast cancer cells was probably mediated by SET8, as ectopic expression of SET8 was able to rescue the effect of miR-7 on tumor migration and invasion (Fig. 3, E and F). Together, these results support an argument that miR-7 suppresses EMT and invasion of breast cancer cells and that this activity is, at least partially, mediated by targeting SET8.

miR-7 Promotes Spontaneous DNA Damages and Sensitizes Cells to Induced DNA Damages—It was reported that loss of function of SET8 was associated with decreases in H4K20me1 and increases in DNA double-strand breaks (15). In light of our observation that SET8 is subjected to negative regulation by
miR-7, we asked the question what role, if any, miR-7 plays in DNA damage response. For this purpose, MCF-7 cells were treated with control or miR-7 mimics for 36 h and challenged with DNA-damaging agents, etoposide (VP-16), or camptothecin for 4 h. Western blotting analysis indicated the levels of phosphorylated H2A.X (γH2AX) increased in miR-7 mimic-treated cells both in the presence and absence of VP-16 or camptothecin (Fig. 4A). Consistently, immunofluorescent microscopy showed that, compared with control, miR-7 mimic-treated cells were enriched with nuclear foci of γH2AX, suggesting that miR-7 promotes the accumulation of spontaneous DNA damages (Fig. 4B). Treatment with VP-16 or camptothecin led to intensified enrichments of γH2AX foci, suggesting that miR-7 sensitizes cells to induced DNA damages (Fig. 4B).

To further consolidate miR-7-induced cellular sensitivity to DNA damage, we tested the effect of miR-7 on cell survival and apoptosis under the exposure of DNA damaging agents. For this purpose, MCF-7 cells treated with control or miR-7 mimics were challenged with VP-16 or camptothecin. Clonogenic assays showed that a more pronounced decrease in the cell viability was detected upon exposure of miR-7-overexpressing MCF-7 cells to DNA damaging agents of increased concentrations (Fig. 4C), which could be partially reversed by reconstitution of SET8. Consistently, transfection of miR-7 mimics in MCF-7 cells resulted in an increased number of cells that underwent apoptosis, which was further exacerbated when cells were exposed to VP-16 or camptothecin (Fig. 4D). Moreover, flow cytometry showed miR-7 overexpression caused a delay of cell cycle progression, revealing that miR-7-induced DNA damage could potentially cause the activation of checkpoint and cell cycle arrest (Fig. 4E). Collectively, these observations suggest that miR-7 promotes cellular susceptibility to genotoxic agents, and the effect is, at least partially, through its targeting of SET8.

miR-7 Impedes both HR and NHEJ of DNA Repair—Given the importance of H4K20 methylation in DNA damage response (26, 27), we next explored how miR-7 influences DNA repair. Immunofluorescent staining showed that miR-7 suppressed the formation of nuclear foci of 53BP1 and RAD51, two key proteins that are recruited at damage sites of chromosomes during DNA repair (28–30), in MCF-7 cells under the exposure of VP-16 or camptothecin (Fig. 5A).

To understand further the role of miR-7 in DNA repair, we investigated the pathway where miR-7 impacts DNA repair, HR, or NHEJ. For this purpose, we utilized two stable cell lines, DR-GFP-U2OS and EJ5-GFP-HEK293, which were designed to

**FIGURE 3.** miR-7 suppresses EMT and invasive potential of breast cancer cells by down-regulating SET8. A, the endogenous expressions of SET8 and miR-7 were measured by Western blotting (left panel) and qPCR (right panel) in MDA-MB-231 and MCF-7 cells. Transcript level of miR-7 was normalized to that of U6. Each bar represents the mean ± S.D. for triplicate experiments. p values were determined by Student’s t test. *p < 0.05, B, the effect of miR-7 on cell morphology in MDA-MB-231 cells. Cells were transfected with miR-7 mimics or miR-7 inhibitors with/without SET8 cDNA lacking the 3′-UTR. The morphological alterations were observed by phase-contrast microscopy. C, the effect of miR-7 on expression of epithelial/mesenchymal markers in MDA-MB-231 cells. Cells were transfected with control or miR-7 mimics and/or SET8 cDNA lacking the 3′-UTR. The immunofluorescence staining of epithelial (E-cadherin and γ-catenin) and mesenchymal (vimentin) markers (green) were visualized by confocal microscopy, and DAPI staining (blue) was included to visualize the cell nucleus. D, cells were transfected with miR-7 mimics or miR-7 inhibitors with/without SET8 cDNA lacking the 3′-UTR. Immunoblotting examination of the epithelial/mesenchymal markers in MDA-MB-231 cells using antibodies against the indicated proteins. E, MDA-MB-231 cells were transfected with miR-7 mimics or miR-7 inhibitors with/without SET8 cDNA lacking the 3′-UTR. Wound healing assays were determined by phase-contrast microscopy at the indicated time points. Each bar represents the mean ± S.D. for triplicate experiments, p values were determined by Student’s t test. *p < 0.05. F, MDA-MB-231 cells were transfected with miR-7 mimics or miR-7 inhibitors with/without SET8 cDNA lacking the 3′-UTR for 48 h. Cells were stained for 18 h before transwell invasion assay were performed using Matrigel transwell filters. The invaded cells were stained and counted. The images represent one field under microscopy in each group. Each bar represents the mean ± S.D. for triplicate experiments, p values were determined by Student’s t test. *p < 0.05. NC, negative control; ln-miR-7, miR-7 inhibitors.
Targeting SET8 by microRNA-7

Epigenetic regulation, by means of DNA methylation, histone modification, chromatin remodeling, and miRNAs, play important roles in a wide range of biological processes, such as X-chromosome inactivation (32), imprinting (33), reprogramming (34), and gene silencing (35). Each of the epigenetic mechanisms does not work alone but forms an essential and intricate network with each other. For example, it has been reported that the promoters in about half of miRNAs are associated with CpG islands that may be repressed by DNA methylation (36). Moreover, a series of histone-modifying enzymes are subject to miRNA modulation. For example, the expression of enhancer of Zeste 2 (EZH2) is inhibited by miR-101 (37), miR-26a (38) and miR-214 (39) in cancer cell lines; miR-137 targets LSD1 and forms a feedback regulatory loop to participate in neural stem cell proliferation and differentiation during neural development (40, 41). We report here that miR-7 is a negative regulator of histone methyltransferase SET8 that inhibits H4K20 monomethylation, aiding the understanding of the complexity of the epigenetic regulatory network.

Thus far, more than 1500 human miRNAs are annotated, which regulate ~30% of the total genomic miRNAs, many of which are associated with a carcinogenic potential (42). It has been reported that miR-7 targets, in addition to BCL-2 (43) and YY1 (11), gene transcripts whose protein products are linked to EGF receptor signaling, including EGF receptor itself (9), RAF-1 (9), p21-activated kinase 1 (44), IRS-1 (45), and insulin receptor substrate-2 (10), to regulate cell proliferation and apoptosis. A recent report showed that miR-7 functions as an anti-metastatic microRNA in gastric cancer by targeting IGFR1 (12). Our current study demonstrated that SET8 is an additional target of miR-7. Significantly, we showed that miR-7 suppressed EMT and invasion of MDA-MB-231 cells, consistent with our previous observation that SET8 and TWIST are functional interdependently in promoting EMT and the invasive potential of breast cancer cells (22).

We demonstrated that miR-7 promoted spontaneous DNA damages and impeded DNA repair, extending its role in DNA damage response. Recent studies indicate that miR-421 and miR-182 are also implicated in DNA damage response and affect cellular sensitivity to DNA damaging agents by targeting ATM (46) or BRCA1 (47), respectively. It is reasonable to

FIGURE 4. miR-7 promotes spontaneous DNA damages and sensitizes cells to induced DNA damages. A, MCF-7 cells were treated with control or miR-7 mimics for 36 h and challenged with 80 nM etoposide (VP-16) or 2 μM camptothecin (CPT) for 4 h. Cellular lysates were analyzed by Western blotting using antibodies against the indicated proteins. B, MCF-7 cells were treated with control or miR-7 mimics for 36 h and challenged with 80 nM VP-16 or 2 μM camptothecin for 4 h. Immunofluorescence staining of γ-H2AX, negative control. C, the effect of miR-7 on cell survival under the exposure of genotoxic agents. MCF-7 cells were transfected with control or miR-7 mimics, and/or SET8 siRNA, and subjected to colony survival assays. Cells were stained with crystal violet and air-dried after 14 days. Numbers of colonies (>30 per colony) were scored by light microscope. Each bar represents the mean ± S.D. for triplicate experiments. p values were determined by Student’s t test. *, p < 0.05. D, the effect of miR-7 on cell apoptosis under the exposure of genotoxic agents. MCF-7 cells were treated with control or miR-7 mimics for 24 h, challenged with 40 nM etoposide (VP-16) or 1 μM camptothecin for 36 h, and stained with annexin V/propidium iodide before flow cytometry. E, the effect of miR-7 on cell cycle progression. MCF-7 cells treated with control or miR-7 mimics were synchronized by double thymidine block and released into the cell cycle for the indicated times. Cells were collected for cell cycle analysis by propidium iodide staining and flow cytometry. Experiments were repeated by three times, and the data from a representative experiment are shown. NC, negative control.

measure HR and NHEJ, respectively (26, 27). The systems utilize GFP as a recombination reporter and I-Scel endonuclease for introduction of double-strand breaks. The repair by HR or NHEJ restores functional GFP expression, and a fluorescent signal can be detected by flow cytometry (31). As shown in Fig. 5, B and C, the efficiency of both HR and NHEJ decreased by ~2-fold in miR-7 mimic-treated cells. Ectopic expression of SET8 partially restored repair potential of the cells. It appears that miR-7 impedes both homologous recombination and non-homologous end joining of DNA repair, and the effect is, at least partially, through its targeting of SET8. Significantly, increased miR-7 levels were also detected in MCF-7 cells exposed to VP-16, camptothecin, or UV (Fig. 5D), as measured by qPCR, suggesting that miR-7 could be induced in DNA damage responses.

DISCUSSION

Epigenetic regulation, by means of DNA methylation, histone modification, chromatin remodeling, and miRNAs, play important roles in a wide range of biological processes, such as X-chromosome inactivation (32), imprinting (33), reprogramming (34), and gene silencing (35). Each of the epigenetic mechanisms does not work alone but forms an essential and intricate network with each other. For example, it has been reported that the promoters in about half of miRNAs are associated with CpG islands that may be repressed by DNA methylation (36). Moreover, a series of histone-modifying enzymes are subject to miRNA modulation. For example, the expression of enhancer of Zeste 2 (EZH2) is inhibited by miR-101 (37), miR-26a (38) and miR-214 (39) in cancer cell lines; miR-137 targets LSD1 and forms a feedback regulatory loop to participate in neural stem cell proliferation and differentiation during neural development (40, 41). We report here that miR-7 is a negative regulator of histone methyltransferase SET8 that inhibits H4K20 monomethylation, aiding the understanding of the complexity of the epigenetic regulatory network.

Thus far, more than 1500 human miRNAs are annotated, which regulate ~30% of the total genomic miRNAs, many of which are associated with a carcinogenic potential (42). It has been reported that miR-7 targets, in addition to BCL-2 (43) and YY1 (11), gene transcripts whose protein products are linked to EGF receptor signaling, including EGF receptor itself (9), RAF-1 (9), p21-activated kinase 1 (44), IRS-1 (45), and insulin receptor substrate-2 (10), to regulate cell proliferation and apoptosis. A recent report showed that miR-7 functions as an anti-metastatic microRNA in gastric cancer by targeting IGFR1 (12). Our current study demonstrated that SET8 is an additional target of miR-7. Significantly, we showed that miR-7 suppressed EMT and invasion of MDA-MB-231 cells, consistent with our previous observation that SET8 and TWIST are functionally interdependent in promoting EMT and the invasive potential of breast cancer cells (22).
Targeting SET8 by microRNA-7

Believe that miRNAs, through regulation of the enrichment of various DNA repair factors at the post-transcriptional level, add another layer to the sophisticated regulatory network of DNA repair system, ensuring the efficiency and accuracy of DNA repair.

H4K20 monomethylation has been associated with nucleosome decondensation and DNA damage foci accessible for recruitment of DNA repair proteins following DNA damage (15). Although other as-yet-unidentified substrates cannot be excluded, H4K20 monomethylation appears to be the major effector of SET8 and a crucial mediator of its impact on DNA repair. In support of this argument, it is noteworthy that embryonic stem cells lacking SET8 display massive DNA damage and γH2AX foci (16). In addition, SET8 is somewhat unique among histone methyltransferases in that its protein level oscillates during cell cycle progression. It is highly expressed during G1/M and early G2, and is absent during S phase (48, 49). Thus, tight regulation of SET8 appears to be essential for proper cell cycle progression. Recent studies showed that SET8 is also regulated at the transcriptional level (50, 51). Our observation that SET8 is targeted by miR-7 revealed a mechanism by which SET8 is regulated post-transcriptionally, underscoring the importance of SET8 regulation.

Finally, despite the progresses made in miRNA research, a comprehensive understanding of the interplay/cross-talk among their multiple targets is urgently needed. With the identification of a broad spectrum of target genes for miRNAs, it is difficult and improper to divide the miRNAs into oncogenes and tumor suppressors due to the diverse activities of target genes and various cellular contexts for a particular miRNA. This is probably also true for miR-7. Nevertheless, our experiments provide a molecular mechanism underlying the regulation of SET8 and extend the biological function of miR-7 to DNA damage response, supporting the pursuit of miR-7 as a potential target for breast cancer intervention.

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