

The lncRNA-GAS5/miR-221-3p/DKK2 Axis Modulates ABCB1-Mediated Adriamycin Resistance of Breast Cancer via the Wnt/ β -Catenin Signaling Pathway

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Drug resistance, including adriamycin (ADR)-based therapeutic resistance, is a crucial cause of chemotherapy failure in breast cancer treatment. Acquired chemoresistance has been identified to be closely associated with the overexpression of P-glycoprotein (P-gp/ABCB1). Long non-coding RNA (lncRNA) growth arrest-specific 5 (GAS5) can be involved in carcinogenesis; however, its roles in ABCB1-mediated ADR resistance are poorly understood. In this study, we identified a panel of differentially expressed lncRNAs, mRNAs, and miRNAs in MCF-7 and MCF-7/ADR cell lines through RNA sequencing (RNA-seq) technologies. GAS5 level was downregulated whereas ABCB1 level was upregulated in the resistant breast cancer tissues and cells. Overexpression of GAS5 significantly enhanced the ADR sensitivity and apoptosis, and it inhibited the efflux function and expression of ABCB1 *in vitro*, while knockdown of GAS5 had the opposite effects. Further mechanism-related investigations indicated that GAS5 acted as an endogenous “sponge” by competing for miR-221-3p binding to regulate its target dickkopf 2 (DKK2), and then it inhibited the activation of the Wnt/ β -catenin pathway. Functionally, GAS5 enhanced the anti-tumor effect of ADR *in vivo*. Collectively, our findings reveal that GAS5 exerted regulatory function in ADR resistance possibly through the miR-221-3p/DKK2 axis, providing a novel approach to develop promising therapeutic strategy for overcoming chemoresistance in breast cancer patients.

many factors and causes of acquired drug resistance, overexpression of P-glycoprotein (P-gp/ABCB1) encoding by the multidrug resistance 1 (*MDR1*) gene plays a significant role in drug resistance due to the increased energy-dependent efflux of cytotoxic drugs from cancer cells.^{5,6} So far, however, the detailed mechanisms of drug resistance in cancer cells are complex and have not been fully elucidated. Therefore, a better understanding of the molecular mechanisms of drug resistance in breast cancer is necessary in order to identify the promising therapeutic targets and improve the efficiency of chemotherapy for breast cancer patients.

Long non-coding RNAs (lncRNAs) are a class of transcripts more than 200 nt in length and limited protein-coding ability, which have been confirmed to be involved in many cellular and genomic processes associated with carcinogenesis and drug resistance/sensitivity.^{3,7} Moreover, the importance of lncRNAs in the resistance of breast cancer to multiple drugs has been discussed. For instance, lncRNA TINCR (terminal differentiation-induced non-coding RNA) enhanced trastuzumab resistance and migration, invasion, and the accompanied epithelial-mesenchymal transition (EMT) process in breast cancer by competitive binding to miR-125b, upregulating HER-2 (human epidermal growth factor receptor-2) and *Snail-1*.⁸ Knockdown of H19 restored chemosensitivity in a paclitaxel-resistant triple-negative breast cancer cell line by regulating the Akt-mediated apoptotic signaling pathway.⁹ Moreover, knockdown of H19 repressed the expression of transcription factors related to the Wnt pathway and EMT, and it improved

INTRODUCTION

Breast cancer is the most common cancer in women and one of the most common causes of cancer death worldwide, accounting for 30% in diagnosed new cases and 15% in estimated deaths.¹ Anthracycline genotoxic drug epirubicin or adriamycin (ADR) is a major chemotherapeutic drug that has been widely applied for breast cancer patients; however, the acquired drug resistance frequently restricted its curative effect and thus led to failure in the clinic.^{2–4} Among

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Table 1. Clinical Characteristics of Breast Cancer Patients

Clinicopathological Data	n	Percentage (%)
Age (years)		
≤ 50	18	69.2
>50	8	30.8
Primary tumor size (cm)		
≤ 5	16	61.5
>5	10	38.5
Lymph node metastasis		
Present	21	80.8
Absent	5	19.2
TNM stage		
I–II	11	42.3
III–IV	15	57.7
Histopathological grade		
I–II	15	57.7
III	11	42.3
ER status		
Positive	13	50.0
Negative	13	50.0
PR status		
Positive	11	42.3
Negative	15	57.7
HER-2 status		
Positive	12	46.2
Negative	14	53.8
Molecular subtype		
Luminal A	5	19.2
Luminal B	9	34.6
Her-2 overexpressing	8	30.8
Triple negative	4	15.4

TNM, tumor, node, metastasis; ER, estrogen receptor; PR, progesterone receptor; HER-2, human epidermal growth factor receptor-2.

tamoxifen sensitivity for promoting cell growth and inhibiting apoptosis in tamoxifen-resistant breast cancer cells.¹⁰ lncRNA SNHG14 contributed to trastuzumab resistance and tumorigenesis in breast cancer cells through regulating *PABPC1* expression by H3K27 acetylation.¹¹ Furthermore, extracellular lncRNA-SNHG14 was able to be incorporated into exosomes and transmitted to sensitive breast cancer cells, thus inducing trastuzumab resistance.¹² lncRNA growth arrest-specific 5 (GAS5) is an lncRNA 650 bases in length, originally isolated from NIH 3T3 cells using subtraction hybridization in 1988,¹³ which was downregulated in breast cancer samples and cells.^{14,15} A recent report demonstrated that upregulation of GAS5 alleviated tamoxifen resistance by acting as a molecular sponge of miR-222, leading to the de-repression of its endogenous target phosphatase and tensin homologs (PTENs) in breast cancer.¹⁶ In addition,

GAS5 can sensitize tumor cells to UV irradiation and doxorubicin, as well as suppress cell invasion by regulating PTENs and PDCD4 through miR-21.¹⁷ Although GAS5 has been suggested to play roles in chemoresistance, the underlying mechanism of GAS5-mediated gene expression having an impact on drug resistance is still elusive.

In this study, we attempted to explore the contributions of GAS5 to the ADR resistance in breast cancer and explore the potential mechanisms. We found that GAS5 expression was decreased whereas ABCB1 was increased in breast cancer-resistant patients and cell lines (Table 1). Furthermore, our results show, for the first time, that GAS5 modulates ABCB1-mediated ADR resistance by targeting the miR-221-3p/dickkopf 2 (DKK2)/β-catenin pathway in breast cancer cells. Thus, GAS5 maybe a promising therapeutic biomarker and target for the treatment of breast cancer patients with ADR resistance.

RESULTS

GAS5 Is Downregulated while ABCB1 Is Upregulated in Breast Cancer Tissues and Cell Lines

The RNA expression profiles were obtained from MCF-7 (human breast cell line) and ADR-resistant MCF-7 cells (MCF-7/ADR) by RNA sequencing (RNA-seq). A total of 3,811 lncRNAs and 4,088 mRNAs were detected with the criteria of absolute false discovery rate (FDR) value <0.05 and p value <0.01. Among them, 1,264 lncRNAs and 1,943 mRNAs were significantly downregulated, whereas 2,547 lncRNAs and 2,145 mRNAs were significantly upregulated in the MCF-7/ADR cells than that in MCF-7 cells (Figures 1A and 1B, all p < 0.01; >3.0-fold). The 10 lncRNAs and mRNAs with the largest fold changes are shown in Tables 2 and 3. Of particular interest, we noticed that GAS5 was one of the most prominently downregulated lncRNAs and ABCB1 was the most upregulated mRNA in the ADR-resistant breast cancer cells. This suggested that low GAS5 expression and high ABCB1 expression were associated with ADR resistance. Next, we examined the expressions of GAS5 and ABCB1 in clinical samples and cells.

According to RECIST criteria, we evaluated the clinical responses to neoadjuvant chemotherapy (NAC) in 26 patients. In our study, a total of 6 patients (23%) had a partial response (PR) and 4 patients (15%) had complete responses (CRs), which were regarded as clinical responders, whereas 10 patients (38%) had stable disease (SD) and 6 patients (23%) had progressive disease (PD), which were regarded as non-responders. Therefore, in our study 10 patients were responders and 16 patients were non-responders. To investigate the potential roles of GAS5 and ABCB1 for response to chemotherapy in breast cancer patients, GAS5 and ABCB1 expression were examined by quantitative real-time PCR in breast cancer tissues of clinical responders (n = 10) and non-responders (n = 16). As shown in Figure 1C, GAS5 expression was considerably downregulated in non-responders compared with that in the responders. On the contrary, ABCB1 expression was upregulated in non-responders compared with that in the responders (Figure 1D). Moreover, the mRNA expression level of ABCB1 was positively correlated with

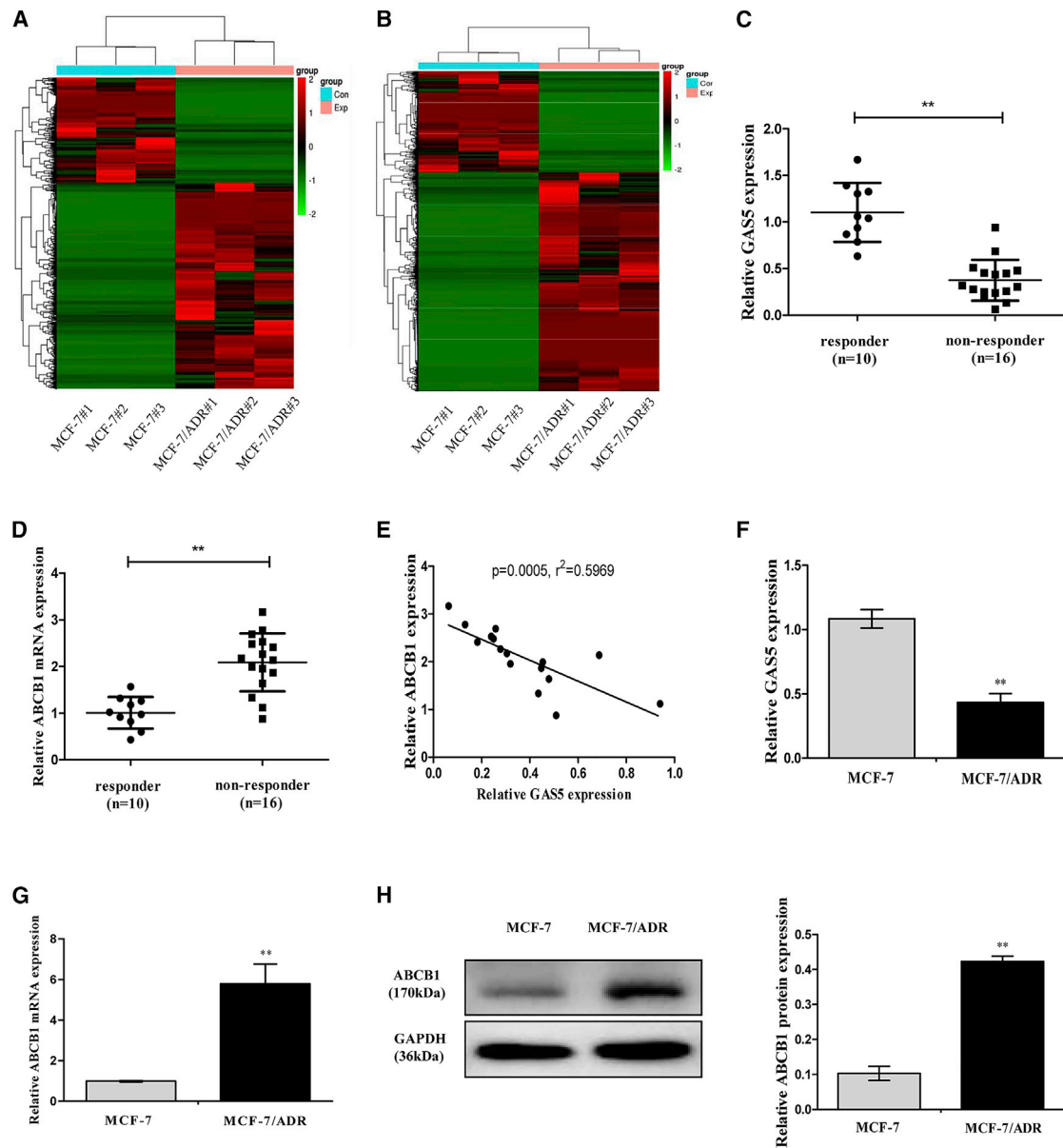


Figure 1. GAS5 and ABCB1 Expression Levels in Breast Cancer Tissues and Cells

(A) Heatmap of differentially expressed lncRNAs in MCF-7 and MCF-7/ADR cells ($n = 3$). (B) Heatmap of differentially expressed mRNAs in MCF-7 and MCF-7/ADR cells ($n = 3$). (C and D) GAS5 expression (C) and ABCB1 mRNA expression (D) in breast cancer tissues from responder and non-responder patients were detected by quantitative real-time PCR. (E) Negative correlation between GAS5 and ABCB1 expressions in breast cancer tissues. (F and G) GAS5 expression (F) and ABCB1 mRNA expression (G) in MCF-7 and MCF-7/ADR cells were detected by quantitative real-time PCR. (H) The ABCB1 protein expression in MCF-7 and MCF-7/ADR cells was detected by western blot analysis. Data are shown as the mean \pm SD ($n = 3$). * $p < 0.05$, ** $p < 0.01$ versus respective control.

GAS5 level in breast cancer tissues (Figure 1E). Due to the relevance between GAS5 and ABCB1 in ADR resistance, we further evaluated the GAS5 and ABCB1 expressions in MCF-7 and MCF-7/ADR cells. The relative expression level of GAS5 was strongly decreased (Figure 1F) whereas ABCB1 was significantly increased (Figures 1G and 1H) by more than 2-fold in the MCF-7/ADR cells compared to the parental MCF-7 cells, indicating that GAS5 and ABCB1 might

play key roles in the development of ADR resistance in breast cancer.

GAS5 Influenced Chemoresistance in Breast Cancer Cells

Afterward, we investigated whether GAS5 is associated with the ADR resistance of breast cancer cells. The 50% inhibition concentration (IC_{50}) of ADR was measured to confirm the ADR resistance of

Table 2. Characteristics of lncRNAs with the Largest Fold Change

lncRNA	Gene ID	Fold Change	Regulation	p Value	FDR
CYTOR	ENSG00000222041.10	5.092170527	up	1.0821E-159	9.3006E-157
C10orf25	ENSG00000165511.6	7.153890466	up	1.904E-132	1.1255E-129
TPRG1-AS1	ENSG00000234076.1	5.672774218	up	7.442E-108	3.1891E-105
ADGRL2	ENSG00000117114.19	5.005623404	up	1.63691E-87	5.04765E-85
LINC00707	ENSG00000238266.1	12.8054899	up	1.71569E-80	4.69952E-78
LINC00886	ENSG00000240875.5	5.154727315	down	5.2902E-173	5.775E-170
AL035661.1	ENSG00000274173.1	8.484605987	down	7.65708E-94	2.6819E-91
GAS5	ENSG00000234741.7	5.038222071	down	1.91332E-89	6.15376E-87
AC010503.4	ENSG00000275234.1	6.181951912	down	5.82009E-76	1.45558E-73
FP325330.3	ENSG00000279712.1	7.88445797	down	2.33947E-70	5.10034E-68

MCF-7/ADR cells in comparison with its parental MCF-7 cells. As shown in Figure 2A, the IC₅₀ value of ADR in MCF-7/ADR cells was almost 160-fold compared to that in MCF-7 cells. To further uncover the function of GAS5 on the sensitivity of ADR, the breast cancer cells were transfected with GAS5 overexpression plasmid (pcDNA-GAS5) or two specific GAS5 small interfering RNAs (siRNAs) (si-GAS5#1 and si-GAS5#2). As expected, GAS5 expression was markedly upregulated in MCF-7/ADR cells when transfected with pcDNA-GAS5, whereas GAS5 expression was down-regulated by 64% by si-GAS5#1 and 41% by si-GAS5#2 compared with their counterparts, respectively (Figure 2B). Additionally, the value of IC₅₀ of ADR and cell proliferation capacity were all strongly decreased in MCF-7/ADR cells by overexpression of GAS5, while introduction with si-GAS5#1 or si-GAS5#2 in MCF-7 cells showed an inverse effect (Figures 2C and 2D). Furthermore, flow cytometry analysis observed that ADR-induced apoptosis was remarkably increased in pcDNA-GAS5-transfected MCF-7/ADR cells and decreased in si-GAS5#1- or si-GAS5#2-transfected MCF-7 cells (Figure 2E). Overall, all of the data concluded that GAS5 reversed the breast cancer cell chemoresistance to ADR-based chemotherapy *in vitro*; here, we further examined the effect of GAS5 on the functions and expressions of ABCB1.

Table 3. Characteristics of mRNAs with the Largest Fold Change

Gene Name	Gene ID	Fold Change	Regulation
ABCB1	ENSG00000085563.14	12.48764005	up
GALC	ENSG00000054983.16	10.31859095	up
CALD1	ENSG00000122786.19	9.684997032	up
COL12A1	ENSG00000111799.20	7.649996364	up
AEBP1	ENSG00000106624.10	6.849429416	up
GRHL2	ENSG00000083307.10	11.78960888	down
SPDEF	ENSG00000124664.10	10.53368044	down
PRLR	ENSG00000113494.16	9.023956295	down
ADAMTS19	ENSG00000145808.9	8.368612293	down
MAP7	ENSG00000135525.18	7.322561697	down

Rhodamine-123 (Rh123), which is an ABCB1-specific fluorescent dye, was used to measure the efflux function of ABCB1.¹⁸ As described in Figure 2F, compared with sensitive MCF-7 cells, lesser accumulation of Rh123 was significantly observed in resistant MCF-7/ADR cells. When the GAS5 expression was overexpressed by pcDNA-GAS5, the Rh123 accumulation in MCF-7/ADR cells was increased 1.78-fold compared to pcDNA-negative control (NC) (empty pcDNA3.1 vector) group cell levels. Consistently, the Rh123 accumulation was decreased almost 1.52-fold in MCF-7 cells transfected with specific si-GAS5#1 than in the NC group siRNA (si-NC), indicating that GAS5 inhibited the cellular efflux function of ABCB1. Furthermore, overexpression of GAS5 dramatically decreased ABCB1 mRNA and protein levels by almost 60% in MCF-7/ADR cells, and knockdown of GAS5 by transferring si-GAS5#1 increased the levels by more than 50% in MCF-7 cells (Figures 2G and 2H). In total, our results exhibited that GAS5 might restrain ABCB1 function and expression, thus influencing the chemoresistance of breast cancer cells.

GAS5 Inhibited miR-221-3p Expression by Direct Interaction

In order to explore the potential mechanism of GAS5 involved in ADR sensitivity in breast cancer cells, the online databases DIANA-LncBase, miRcode, and RNA-seq were applied to predict and confirm the most underlying microRNA (miRNA) that can be interacted with GAS5. We selected and identified the 456 miRNAs that were significantly differentially expressed by RNA-seq (Figure 3A, all $p < 0.01$; >3.0 -fold). Of all of the 456 miRNAs, we found that miR-221-3p was the most upregulated miRNA in the MCF-7/ADR cells (Table 4). Interestingly, the results of online bioinformatics analysis indicated that complementary sites exist between miR-221-3p and GAS5 (Figure 3B). Moreover, the subcellular location of GAS5 was determined by quantitative real-time PCR. The results showed that GAS5 was mainly localized in the cytoplasmic fraction, indicating that GAS5 had a chance to interact with cytoplasmic miRNAs (Figure 3C). The further dual-luciferase reporter assay and RNA immunoprecipitation (RIP) assay validated the direct binding between miR-221-3p and GAS5 in MCF-7/ADR cells. In the dual-luciferase reporter assay, co-transfection of MCF-7/ADR cells with GAS5

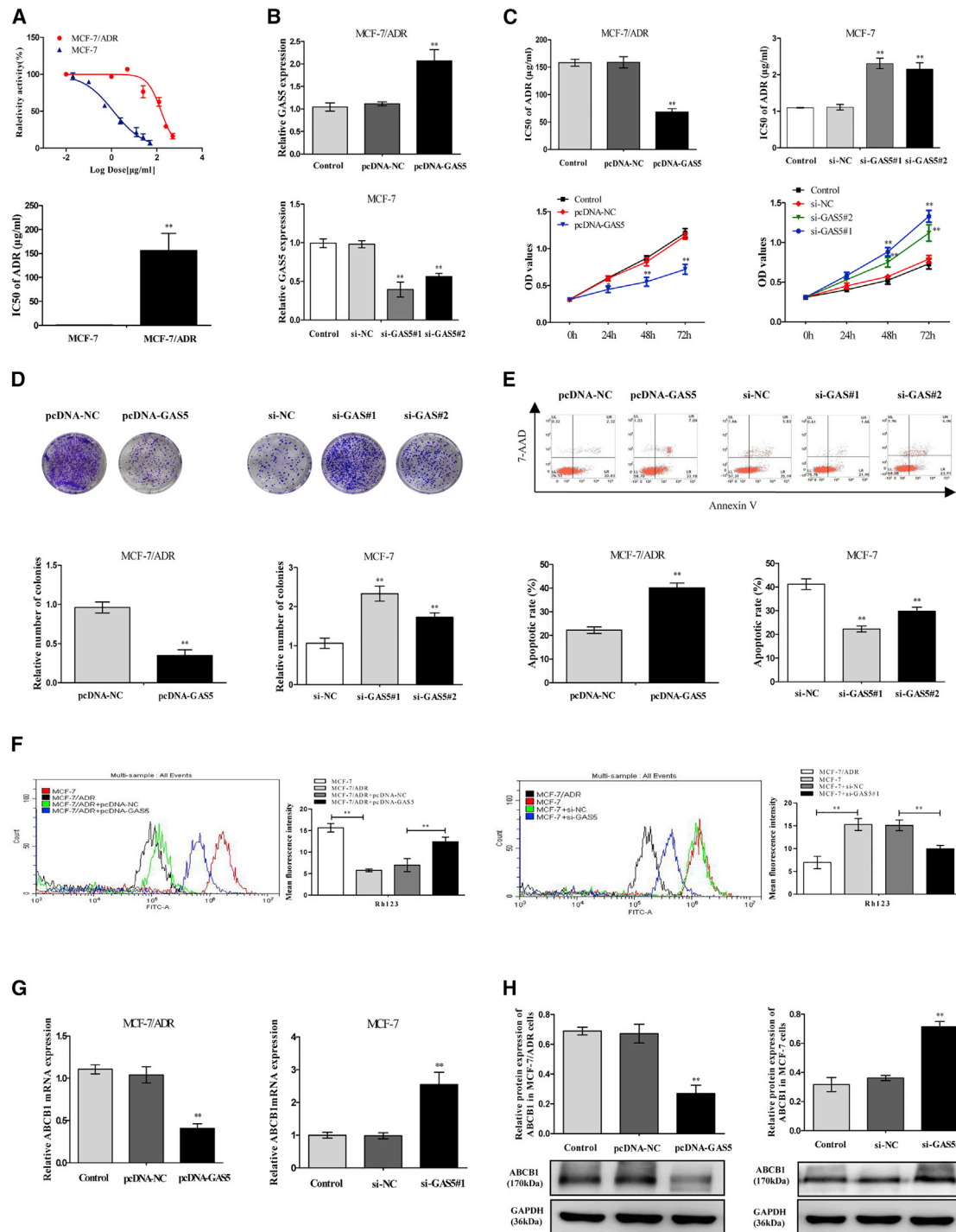


Figure 2. GAS5 Was Associated with ADR Resistance of Breast Cancer Cells

(A) Cells were exposed to a series dose of ADR (0.02, 0.1, 0.5, 2.5, 12.5, and 25 $\mu\text{g/ml}$ for MCF-7, and 0.5, 1, 5, 25, 125, 250 $\mu\text{g/ml}$ for MCF-7/ADR) for 48 h; the cell viability of cells and the 50% inhibitory concentration (IC₅₀) of ADR were then detected using a CCK-8 assay. (B) MCF-7/ADR cells were transfected with pcDNA-GAS5, and MCF-7 cells were introduced with two synthesized individual GAS5 siRNAs (si-GAS5#1 and si-GAS5#2). The GAS5 expression was detected by quantitative real-time PCR. (C) pcDNA-GAS5-transfected MCF-7/ADR cells and si-GAS5#1- or si-GAS5#2-transfected MCF-7 cells were treated with various concentrations of ADR for 48 h, and IC₅₀s of

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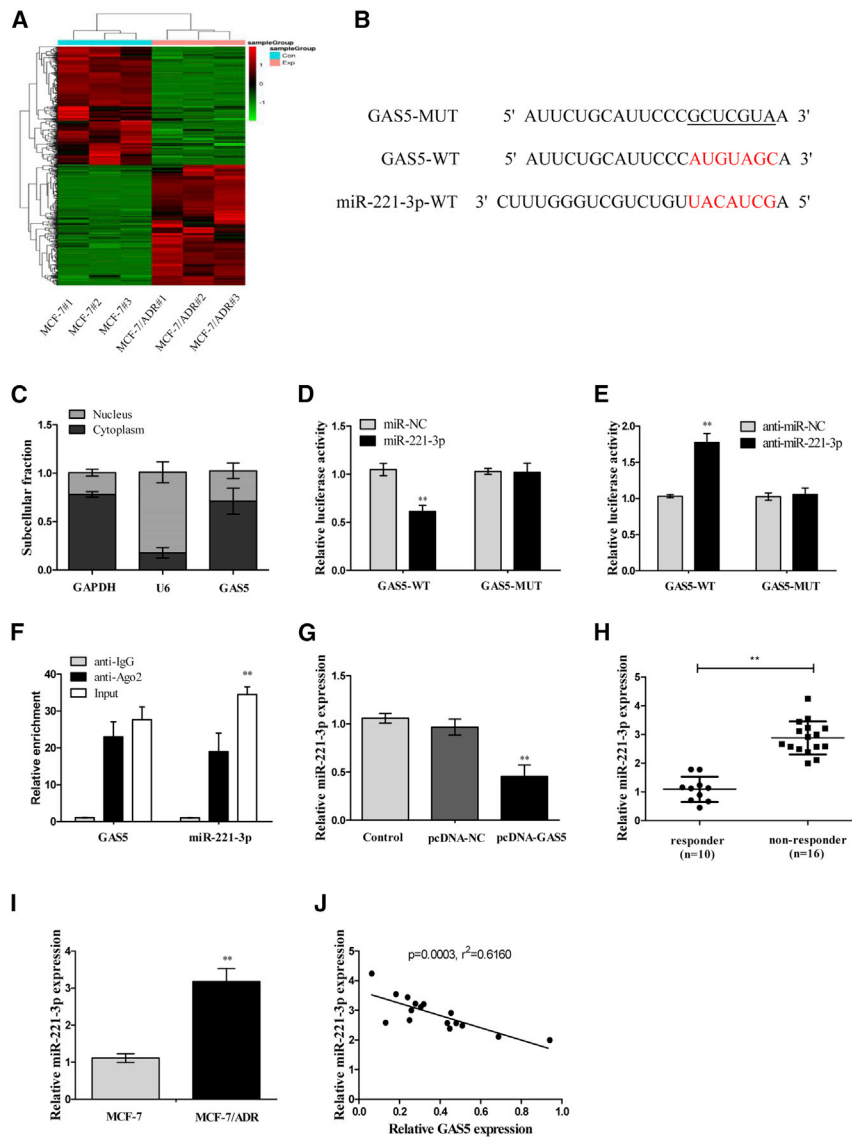


Figure 3. GAS5 Directly Suppressed miR-221-3p Expression in Breast Cancer Cells

(A) The differently expressed miRNAs between MCF-7 and MCF-7/ADR cells were elucidated by a heatmap (n = 3). (B) Sequence alignment of miR-221-3p with the putative binding sites within the wild-type regions of GAS5. (C) The subcellular location of GAS5 in MCF-7 cells was identified by a subcellular fractionation assay. GAPDH and U6 were used as internal references. (D) Luciferase activity in MCF-7/ADR cells co-transfected with the wild-type or mutant GAS5 reporters (GAS5-WT or GAS5-MUT) and miR-221-3p mimics or miR-control (Con). (E) Luciferase activity in MCF-7/ADR cells co-transfected with GAS5-WT or GAS5-MUT and anti-miR-221-3p or anti-miR-Con. (F) The GAS5 and miR-221-3p levels isolated from Ago2 and IgG immunoprecipitates derived from MCF-7/ADR cells were examined by quantitative real-time PCR. (G) The miR-221-3p expression in MCF-7/ADR cells transfected with pcDNA-GAS5 was examined by quantitative real-time PCR. (H) The miR-221-3p expression in breast cancer tissues was examined by quantitative real-time PCR. (I) The miR-221-3p expression in MCF-7 and MCF-7/ADR cells was examined by quantitative real-time PCR. (J) Pearson's analysis shows a negative correlation between GAS5 and miR-221-3p expressions in breast cancer tissues. Data are shown as the mean ± SD (n = 3). *p < 0.05, **p < 0.01 versus respective control.

effect of GAS5 on miR-221-3p expression was detected in MCF-7/ADR cells transfected with pcDNA-GAS5. As shown in Figure 3G, miR-221-3p was downregulated by ectopic GAS5. Subsequently, we detected the expression of miR-221-3p in breast tumor tissues and cells. It was shown that miR-221-3p was strongly significantly upregulated in non-responders compared with that in the responders (Figure 3H), and miR-221-3p expression in MCF-7/ADR cells was about 3-fold greater than that in corresponding controls (Figure 3I). Additionally, Pearson's correlation analysis demonstrated that miR-221-3p expression was inversely correlated with GAS5 expression in breast tumor tissues (Figure 3J). All of these results suggested that GAS5 could directly bind to miR-221-3p and regulate its expression.

miR-221-3p Mediates the Effects of GAS5 on the ADR Resistance of Breast Cancer Cells

Next, we attempted to explore whether miR-221-3p mediated the effect of GAS5 on the chemoresistance of breast cancer cells. The results indicated that GAS5-triggered reduction of miR-221-3p levels was

wild-type (WT) reporter vector (GAS5-WT) and miR-221-3p mimics remarkably decreased the luciferase activity of GAS5 compared to those transfected with GAS5 mutant (MUT) type reporter vector (GAS5-MUT) (Figure 3D), and downregulation of miR-221-3p by introducing anti-miR-221-3p clearly promoted the luciferase activity (Figure 3E). However, the mutant reporter activities in all transfected cells had no evident effects (Figures 3D and 3E). Moreover, RIP data revealed that GAS5 and miR-221-3p were highly enriched in the complex precipitated by Argonaute2 (Ago2) antibody relative to nonspecific immunoglobulin G (IgG) control antibody (Figure 3F). Then, the

ADR and cell proliferation capacity were measured by a CCK-8 assay. (D) The colony-forming ability was measured by a colony-forming assay. (E) Cell apoptosis was evaluated by flow cytometry. (F) Representative fluorescence-activated cell sorting (FACS) histograms of Rh123 accumulation and intensity of Rh123 fluorescence in pcDNA-GAS5-transfected MCF-7/ADR cells and in si-GAS5#1-transfected MCF-7 cells. (G) The ABCB1 mRNA expression was examined by quantitative real-time PCR. (H) The ABCB1 protein expression was examined by western blot analysis. Data are shown as the mean ± SD (n = 3). *p < 0.05, **p < 0.01 versus respective control.

Table 4. Characteristics of miRNAs with the Largest Fold Change

miRNA	Chromosome	Fold Change	Regulation	p Value	FDR
hsa-miR-221-3p	chrX	7.287296594	up	1.5134E-250	2.8823E-248
hsa-miR-222-3p	chrX	7.285851132	up	1.6712E-248	2.8644E-246
hsa-miR-363-3p	chrX	6.407126736	up	9.8099E-243	1.5286E-240
hsa-miR-29a-3p	chr7	5.79567643	up	6.2988E-232	8.9968E-230
hsa-miR-10a-5p	chr17	5.595258884	up	1.6657E-228	2.0393E-226
hsa-miR-125b-5p	chr11;chr21	3.515215771	up	7.7306E-227	8.8335E-225
hsa-miR-100-5p	chr11	4.749079454	up	2.4896E-216	2.667E-214
hsa-miR-489-3p	chr7	6.667962520	down	3.8488E-231	5.0746E-229
hsa-miR-200b-3p	chr1	3.960448865	down	8.5356E-209	8.6059E-207
hsa-miR-200c-3p	chr12	6.365063854	down	1.3053E-203	1.2429E-201

significantly reversed by miR-221-3p overexpression in MCF-7/ADR cells (Figure 4A). Consistently, si-GAS5#1-induced promotion of miR-221-3p expression was markedly recovered after depletion of miR-221-3p in MCF-7 cells in comparison to homologous controls (Figure 4A). In the Cell Counting Kit-8 (CCK-8) and colony formation assays, GAS5 decreased the IC₅₀ of ADR and cell proliferation, which were strikingly reversed by introduction with miR-221-3p mimics in MCF-7/ADR cells, whereas si-GAS5#1 showed the opposite effects by introduction with anti-miR-221-3p (Figures 4B and 4C). The data from flow cytometry indicated that GAS5-elicited apoptosis was significantly lowered after upregulation of miR-221-3p in MCF-7/ADR cells, while si-GAS5#1-induced reduction on apoptosis was greatly improved following downregulation of miR-221-3p in MCF-7 cells (Figure 4D). Collectively, these findings demonstrated that GAS5 increased the sensitivity of breast cancer cells to ADR through repressing miR-221-3p.

miR-221-3p Directly Targets DKK2, Regulating ABCB1 Expression

Bioinformatics databases (TargetScan, starBase, and miRDB) analysis revealed that DKK2, which is a critical receptor for activation of the Wnt/ β -catenin signaling pathway, was identified as a potential target of miR-221-3p (Figure 5A). In the meantime, it has been reported that DKK2 is a target of miR-221 in osteosarcoma.¹⁹ Following the above findings, we attempted to explore whether DKK2 was a target of miR-221-3p in breast cancer cells. Later, a luciferase reporter assay was performed to confirm the binding site of miR-221-3p. It indicated that miR-221-3p negatively regulated the luciferase activity of DKK2-WT-3' UTR, rather than that of DKK2-MUT-3' UTR (Figure 5B). Subsequently, DKK2 was distinctly downregulated in non-responders compared with responders with respect to breast cancer tissues (Figure 5C), and DKK2 mRNA expression was decreased more in ADR-resistant MCF-7/ADR cells than that in the MCF-7 control group (Figure 5D). Furthermore, Figures 5E and 5F suggested that mRNA and protein expression levels of DKK2 in MCF-7/ADR cells were inactively regulated by miR-221-3p, while the mRNA and protein expression levels of ABCB1 displayed the contrary effect. Therefore, we concluded

that miR-221-3p directly targets DKK2 and induces ABCB1 in breast cancer cells.

GAS5 Inhibits ABCB1 Expression by Activating the Wnt/ β -Catenin Signaling Pathway through the GAS5/miR-221-3p/DKK2 Axis

To determine whether specific crosstalk existed between GAS5 and DKK2 through competition for miR-221-3p binding (Figure 6A), we conducted luciferase report assays by transfecting DKK2 WT reporter vector (DKK2-WT) into cells together with miR-221-3p mimics or miR-221-3p mimics+pcDNA-GAS5. These data indicated that the fluorescence intensity of the DKK2-WT vector was remarkably inhibited by co-transfection with miR-221-3p mimics compared to the corresponding controls. Additionally, miR-221-3p triggered a decrease in luciferase activity of the DKK2-WT vector, which was substantially counteracted by GAS5 upregulation (Figure 6B). In addition, we found that GAS5 increased the DKK2 expression and decreased the expression of the Wnt/ β -catenin pathway downstream targets c-Myc and cyclin D1 (Figures 6C and 6D). Then, we attempted to explore whether GAS5 regulated ABCB1 expression via activating the Wnt/ β -catenin signaling pathway by the GAS5/miR-221-3p/DKK2 axis. Real-time PCR for rescue experiments revealed that GAS5 positively regulated DKK2 mRNA expression, which was mediated by miR-221-3p. It showed that pcDNA-GAS5 not only increased the DKK2 expression but also decreased the β -catenin, c-Myc, cyclin D1, and ABCB1 expressions, as previously mentioned, while when combined with miR-221-3p mimics or si-DKK2 it reversed the effects of pcDNA-GAS5 (Figures 6E-6G). When combining the preceding findings, the data demonstrated that GAS5 inhibits ABCB1 expression by activating the Wnt/ β -catenin signaling pathway through the GAS5/miR-221-3p/DKK2 axis.

GAS5 Enhances ADR Sensitivity in Breast Cancer *In Vivo*

To further verify the underlying effects and mechanisms of GAS5 in increasing the sensitivity of breast cancer MCF-7/ADR cells to ADR *in vivo*, cells infected with pcDNA-GAS5 or pcDNA-NC were subcutaneously injected into each nude mouse to generate tumor xenograft,

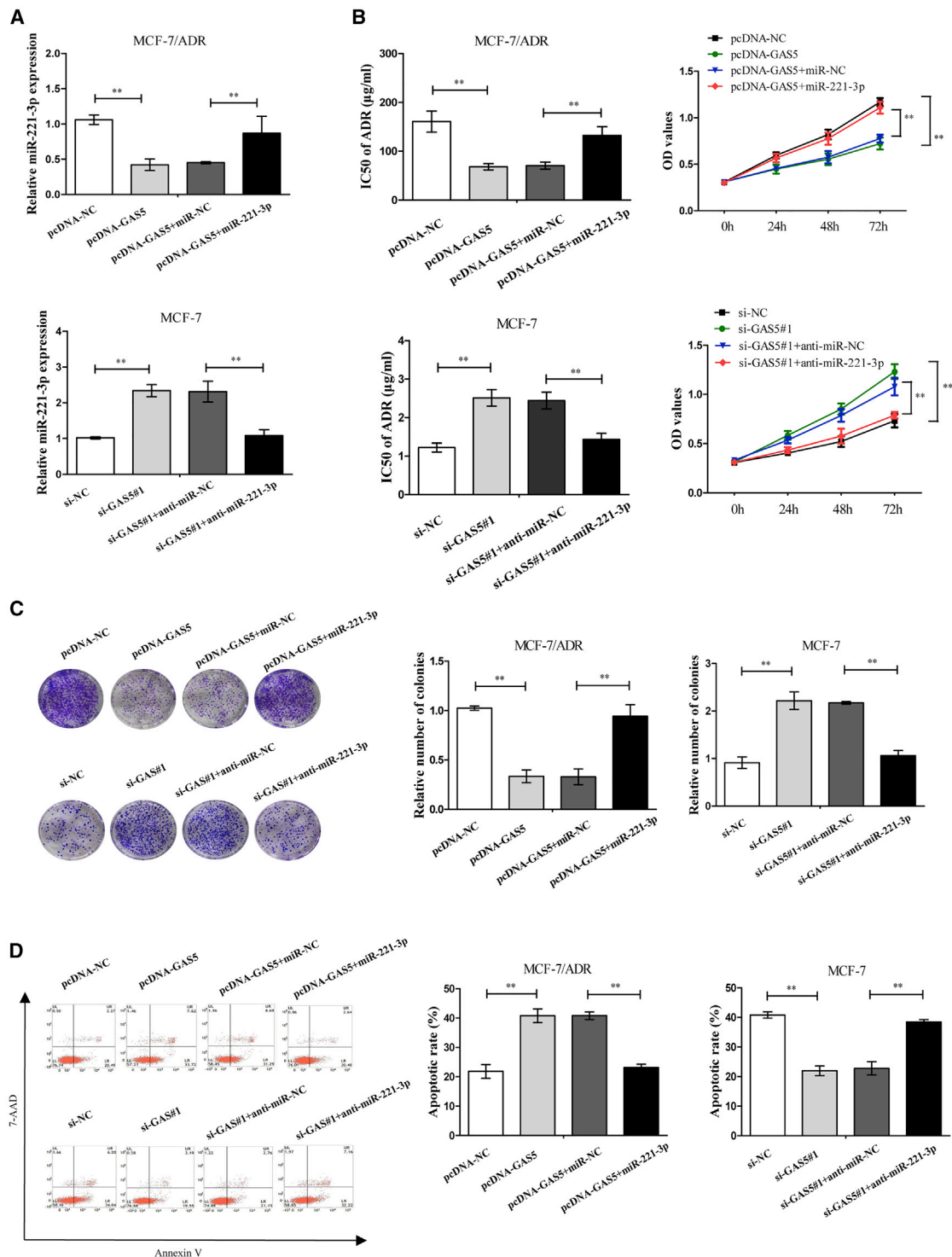


Figure 4. GAS5-Induced ADR Sensitivity of Breast Cancer Cells Was Abated by miR-221-3p Upregulation

MCF-7/ADR cells were transfected with pcDNA-GAS5 alone or together with miR-221-3p mimics, and MCF-7 cells were transfected with si-GAS5#1 alone or together with anti-miR-221-3p. (A) The miR-221-3p expression in MCF-7/ADR and MCF-7 cells was examined by quantitative real-time PCR. (B) IC₅₀ of ADR and cell proliferation capacity were measured by a CCK-8 assay in MCF-7/ADR and MCF-7 cells. (C) The colony-forming ability was measured by a colony-forming assay. (D) Cell apoptosis was evaluated by flow cytometry. The data are shown as the mean ± SD (n = 3). *p < 0.05, **p < 0.01 versus respective control.

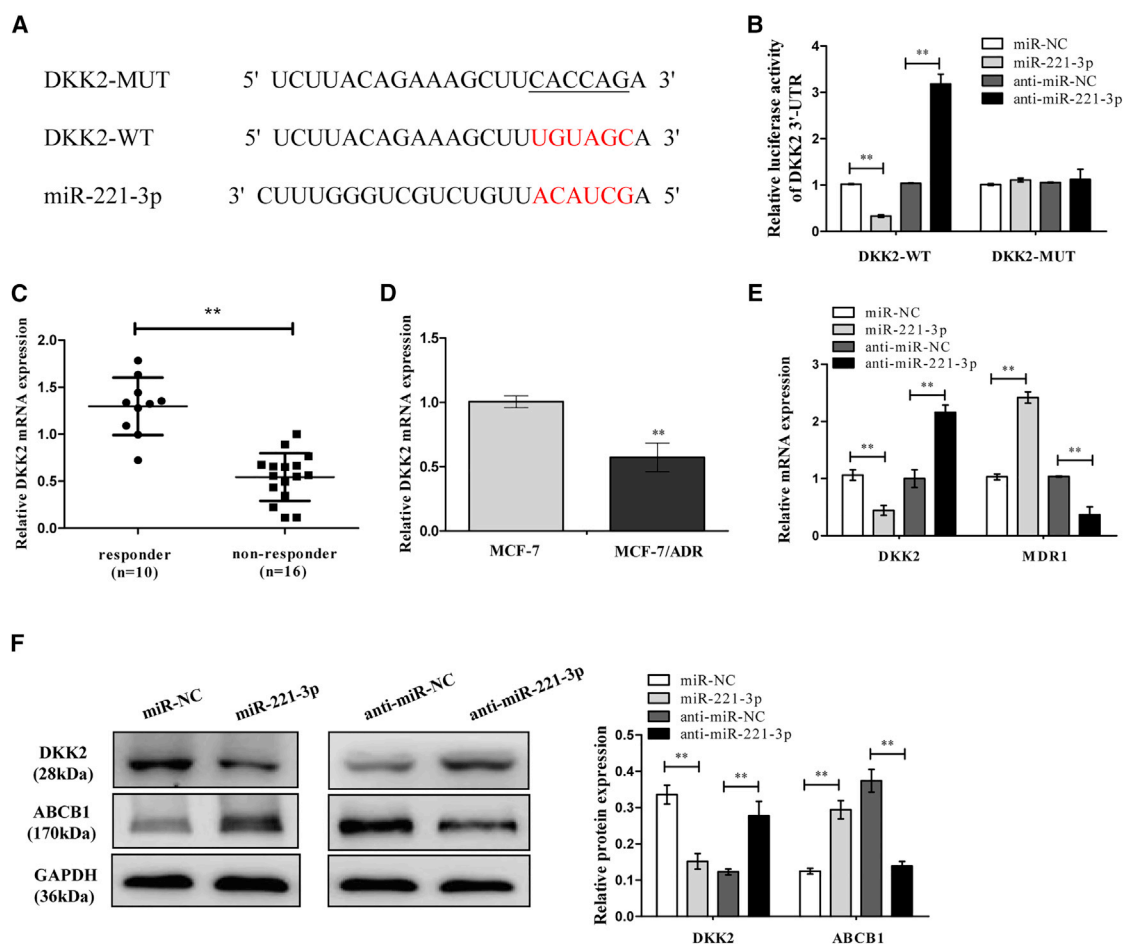


Figure 5. miR-221-3p Targets the 3' UTR of DKK2 and Regulates the ABCB1 Expression

(A) Putative binding sites between DKK2 and miR-221-3p predicted by the TargetScan online website together with the mutant sites of the DKK2-3' UTR region in the MUT-DKK2-3' UTR reporter vector. (B) The luciferase activity was detected in MCF-7/ADR cells transfected with DKK2-WT or DKK2-MUT and miR-221-3p mimics or anti-miR-221-3p. (C) The DKK2 mRNA expression in breast cancer tissues was examined by quantitative real-time PCR. (D) The DKK2 mRNA expression in MCF-7 and MCF-7/ADR cells was examined by quantitative real-time PCR. (E) The MCF-7/ADR cells were transfected with miR-221-3p mimics or anti-miR-221-3p. The DKK2 and ABCB1 mRNA expression levels were examined by quantitative real-time PCR. (F) The DKK2 and ABCB1 protein expression levels were examined by western blot analysis. Data are shown as the mean \pm SD ($n = 3$). * $p < 0.05$, ** $p < 0.01$ versus respective control.

followed by administration with ADR or PBS. As shown in Figures 7A and 7B, GAS5 overexpression or ADR treatment dramatically decreased the tumor volume and weight compared with the control group. Furthermore, a more distinct inhibition on tumor growth was caused by simultaneous GAS5 overexpression together with ADR treatment, suggesting the vital role of GAS5 on ADR sensitivity *in vivo* (Figures 7A and 7B). Consistent with previous findings, we found that the expressions of GAS5 (Figure 7C) and DKK2 (Figure 7E) were increased, but the expressions of miR-221-3p (Figure 7D) and ABCB1 (Figure 7E) were decreased, in tumor tissues derived from GAS5-overexpressed MCF-7/ADR cells with or without ADR treatment. A Western blot assay confirmed that the DKK2 expression level was clearly increased while the ABCB1 level was decreased followed by GAS5 overexpression in tumor xenografts with or without ADR treatment (Figure 7F). Overall, these results

suggested that overexpression of GAS5 enhances ADR sensitivity in breast cancer *in vivo*.

DISCUSSION

During the past decades, the overall prognosis of breast cancer patients diagnosed at the early stage has been remarkably improved through standard chemotherapy after surgical resection. Largely due to drug resistance, however, the 5-year survival rates of breast cancer patients remain dismal.^{20–22} It has been proposed that drug resistance has frequently been associated with elevated expression levels of ABCB1 in breast cancer cells.^{6,23} Overcoming resistance mediated by ABCB1 is still a challenge in breast cancer chemotherapy. Recently, emerging evidence has demonstrated that aberrant lncRNA expression is strongly implicated in the development of drug resistance in breast cancer; however, the molecular mechanisms for

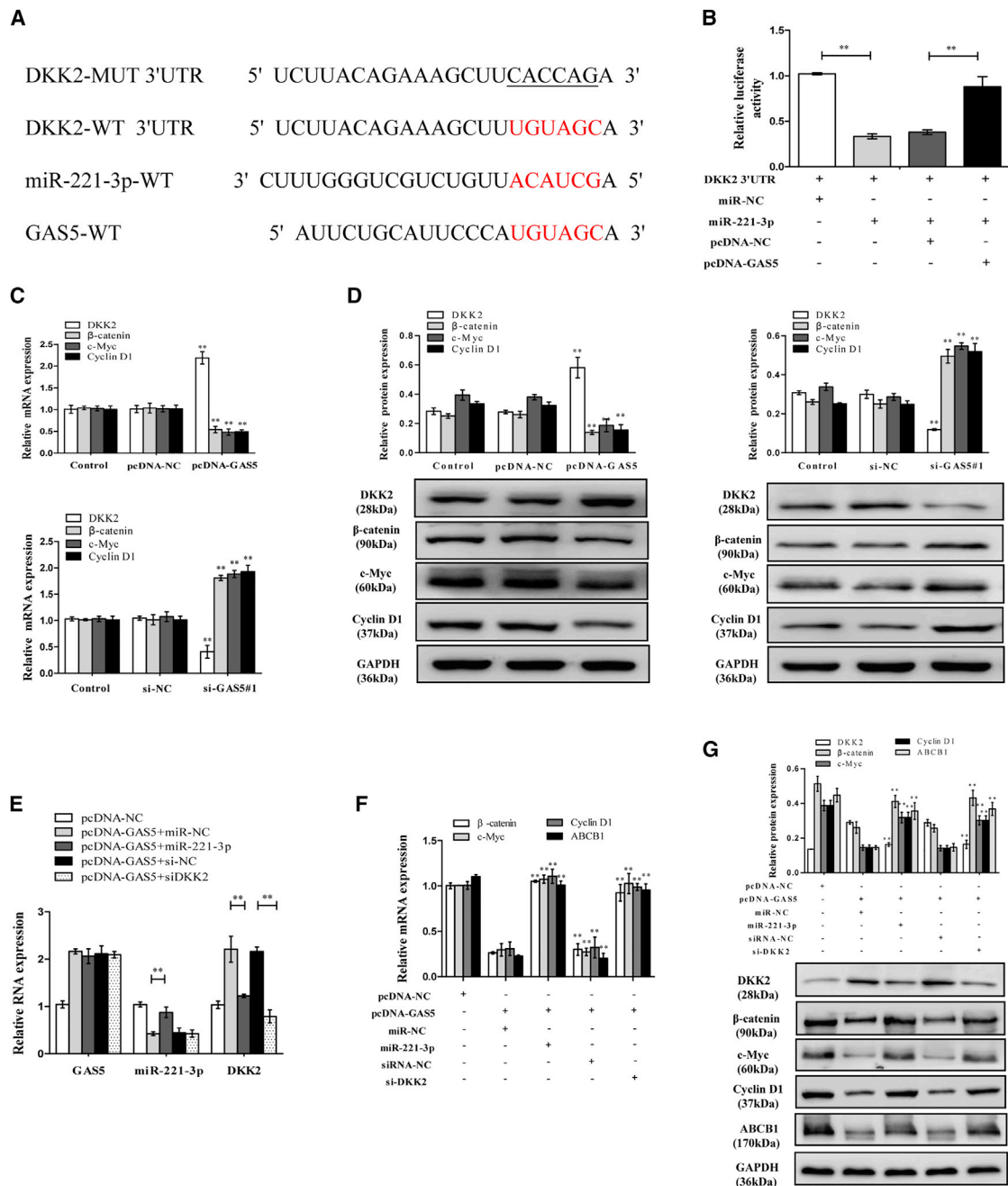


Figure 6. GAS5 Inhibits ABCB1 Expression by Activating the Wnt/ β -Catenin Signaling Pathway through the GAS5/miR-221-3p/DKK2 Axis

(A) Putative miR-221-3p binding sites on GAS5 and DKK2. (B) The dual-luciferase reporter assay was performed by transfecting DKK2-WT vector into MCF-7/ADR cells together with miR-221-3p mimics or miR-221-3p mimics+pcDNA-GAS5. (C) The DKK2, β -catenin, c-Myc, and cyclin D1 mRNA expressions in MCF-7/ADR cells transfected with pcDNA-NC or pcDNA-GAS5 and in MCF-7 cells transfected with si-NC or si-GAS5#1 were detected by quantitative real-time PCR. (D) The DKK2, β -catenin, c-Myc, and cyclin D1 protein expressions in MCF-7/ADR cells transfected with pcDNA-NC or pcDNA-GAS5 and in MCF-7 cells transfected with si-NC or si-GAS5#1 were detected by western blot analysis. (E) MCF-7/ADR cells were transfected with pcDNA-GAS5+miR-NC, pcDNA-GAS5+miR-221-3p mimics, pcDNA-GAS5+siRNA-NC, or pcDNA-GAS5+si-DKK2. The GAS5, miR-221-3p, and DKK2 expressions were examined by quantitative real-time PCR. (F) The DKK2, β -catenin, c-Myc, and cyclin D1 mRNA expressions were examined by quantitative real-time PCR. (G) The DKK2, β -catenin, c-Myc, and cyclin D1 protein expressions were examined by western blot analysis. Data are shown as the mean \pm SD (n = 3). *p < 0.05, **p < 0.01 versus respective control.

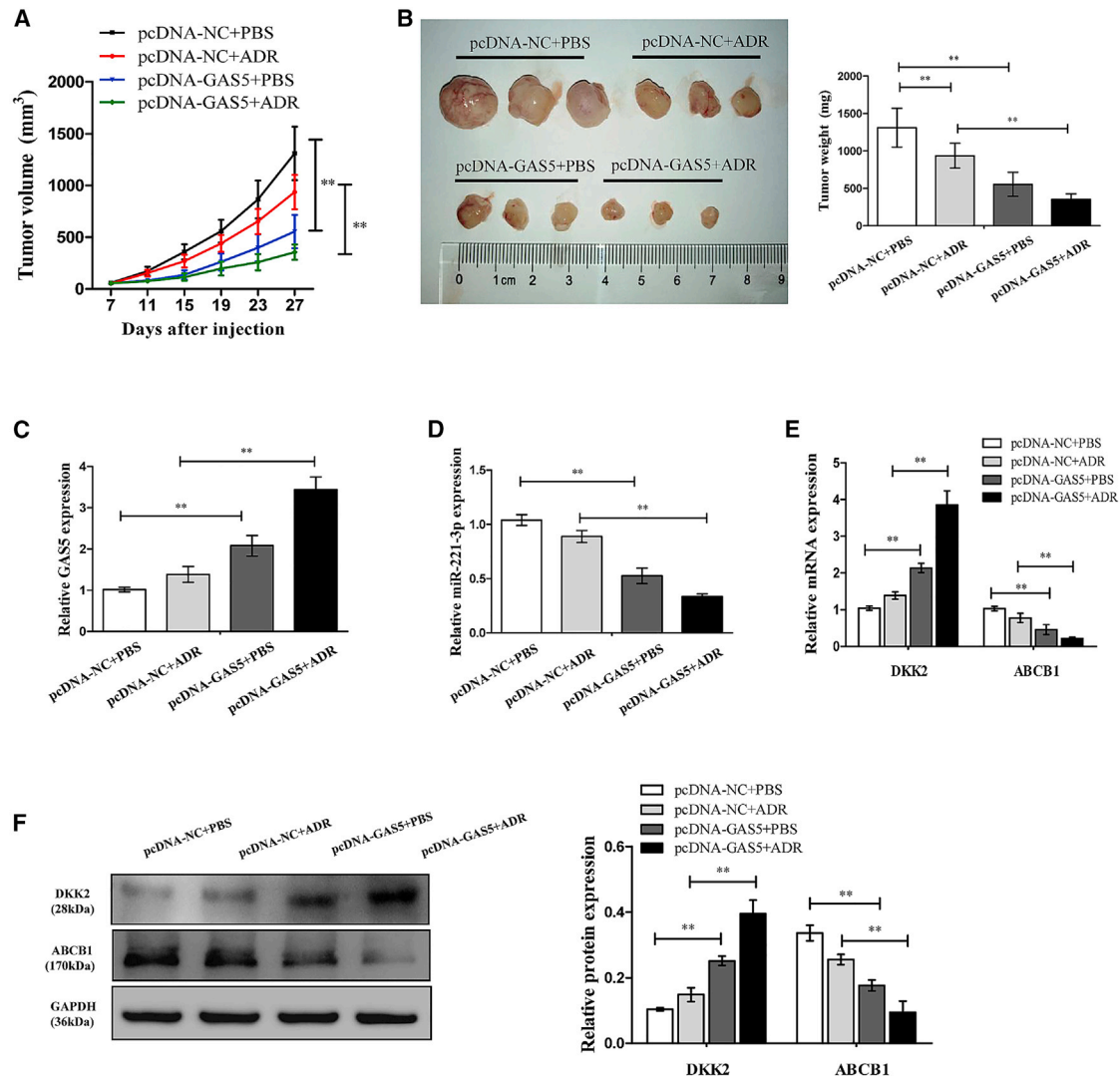


Figure 7. Overexpression of GAS5-Sensitized MCF-7/ADR Cells to ADR *In Vivo*

MCF-7/ADR cells stably infected pcDNA-GAS5 or pcDNA-NC were subcutaneously injected into nude mice, followed by administration with ADR (4 mg/kg) every 4 days from the 7th day after cell inoculation. (A) Tumor volumes were examined every 4 days for 27 days. (B) Representative photographs and average weights of dissected tumors. (C) The GAS5 expression in resected tumor masses was detected by quantitative real-time PCR. (D) The miR-221-3p expression in resected tumor masses was detected by quantitative real-time PCR. (E) The mRNA expression levels of DKK2 and ABCB1 in resected tumor masses were detected by quantitative real-time PCR. (F) The protein expression levels of DKK2 and ABCB1 in resected tumor masses were detected by western blot analysis. Data are shown as the mean \pm SD ($n = 3$). * $p < 0.05$, ** $p < 0.01$ versus respective control.

regulation of ABCB1 by lncRNA in drug resistance have not been well elucidated.

In our present study, based on the analysis results of RNA-seq, we focused on lncRNA GAS5, which significantly contributes to various aspects of cancer biology and has been identified as a critical player of drug resistance in cancer therapy. Liu et al.²⁴ found that GAS5 was downregulated in both pancreatic cancer tissues and cell lines. Overexpression of GAS5 suppressed cell proliferation, cell invasion, and migration and promoted gemcitabine-induced apoptosis by regu-

lating the miR-221/SOCS3 pathway, mediating EMT and tumor stem cell self-renewal. Similarly, GAS5 upregulation could decrease the malignancy and cisplatin resistance of cervical cancer HeLa cells via the miR-21/STAT3 signaling pathway.²⁵ Furthermore, GAS5 improved the development of multidrug resistance of pancreatic cancer cells by regulating the miR-181c-5p/ABCB1/Hippo pathway. A recent study indicated that GAS5 was significantly downregulated in tamoxifen-resistant MCF-7R cells, and overexpression of GAS5 could enhance the cell sensitivity to tamoxifen in MCF-7R cells both *in vitro* and *in vivo*.¹⁶ However, it is still unclear whether GAS5 could

also affect ADR resistance in breast cancer cells. ADR is an ABCB1 substrate, which limits its effectiveness for current use in treating breast cancer patients.²⁶ Based on the above, we sought to determine the potential mechanisms between GAS5 and ABCB1-mediated ADR resistance in breast cancer. Our results found an inverse correlation of expression patterns between GAS5 and ABCB1. The expression level of GAS5 was significantly low in non-responder breast cancer tissues and in MCF-7/ADR cells, whereas expression level of ABCB1 was very high. Overexpression of GAS5 greatly increased the sensitivity and apoptosis of MCF-7/ADR cells to ADR. Conversely, ADR sensitivity and apoptosis were reduced by treatment with si-GAS5. Moreover, we assessed the impact of GAS5 on the efflux function of ABCB1 by Rh123 transport assay. GAS5 upregulation distinctly increased the intracellular accumulation of the substrate drug Rh123, whereas downregulation decreased compared to the cells transfected with NC. We further found that GAS5 upregulation completely suppressed ABCB1 expression. These findings highlight the importance of GAS5 dysfunction in the development of drug resistance.

Considerable research has shown that one of lncRNA's functional mechanisms is the competing endogenous RNA (ceRNA)-miRNA-mRNA regulatory network, which maintains the activity and functional balance of gene circuitry in a cancer cell. Any disturbance of this system may trigger a series of cellular pathological processes. Hence, we used RNA-seq and online software (e.g., DIANA-LncBase, miRcode) to search for the potential target miRNAs of GAS5. miR-221-3p is regarded as a tumor suppressor in human cancers, including breast cancer.^{27,28} Interestingly, miR-221-3p is the top target miRNA when using RNA-seq analysis. Additionally, several conserved cognate sites between GAS5 and miR-221-3p existed, predicting that GAS5 might serve as a ceRNA of miR-221-3p. Li et al.²⁹ revealed that the ceRNA network normally occurs in the cytoplasm. Our study verified that GAS5 was substantially enriched in the cytoplasmic fraction, showing that GAS5 had a chance to interact with cytoplasm miRNAs. Additionally, our data also indicated that the negative correlation between GAS5 and miR-221-3p was confirmed in ADR-resistant breast cancer tissues. In the subsequent luciferase reporter experiments, the results of RIP and quantitative real-time PCR assays indicated that GAS5 could suppress miR-221-3p expression through direct interaction. GAS5 overexpression could suppress the expression of miR-221-3p. Moreover, induced GAS5 increased the sensitivity and apoptosis of MCF-7/ADR cells to ADR, which was antagonized following the restoration of miR-221-3p expression. Knockdown of GAS5 by si-GAS5#1 decreased the sensitivity and apoptosis of MCF-7 cells to ADR, which was fortified after transfection with miR-221-3p inhibitors. All of these data provided evidence that GAS5 sensitized breast cancer cells to ADR via restraining miR-221-3p expression.

Next, we investigated the potential targets of miR-221-3p, and DKK2 was identified as a functional target in ADR-resistant breast cancer MCF-7/ADR cells. DKK2 is a member of the dickkopf family, which is identified as a secreted modulator of Wnt- β -catenin signaling by binding to the lipoprotein receptor-related protein 5/6 (LRP5/6) component of the Wnt receptor complex.³⁰ DKK2 has been reported

to be involved in drug resistance and promote tumor cell survival, proliferation, metastasis, and invasion.³⁰⁻³² Remarkably, previous research found that DKK2 could stimulate tumor metastasis and angiogenesis via accelerating the aerobic glycolysis of human colorectal cancer cells, through a novel VEGF-independent, but energy metabolism-related pathway.³⁰ Wang et al.³¹ showed that miR-221 functions as an oncomiR and controls 5-fluorouracil (5-FU) resistance in esophageal adenocarcinoma cancer by direct targeting of DKK2 expression via modulation of the Wnt/ β -catenin-EMT pathway. In our study, DKK2 expression was decreased in non-responder breast cancer tissues and in MCF-7/ADR cells. In addition, GAS5 upregulation increased DKK2 expression whereas GAS5 knockdown repressed DKK2 expression. Additionally, GAS5 inhibited activation of the Wnt/ β -catenin pathway. Moreover, the correlation between GAS5 and the miR-221-3p/DKK2 axis was determined by a dual-luciferase reporter assay. In the subsequent rescue experiments, we showed that GAS5 upregulation increased the expression of DKK2 and decreased the expressions of ABCB1 and the Wnt/ β -catenin pathway, while these effects were attenuated following miR-221-3p overexpression or DKK2 inhibition. All of these results implied that GAS5 could reverse the ADR resistance by the miR-221-3p/DKK2 axis via the Wnt/ β -catenin pathway in ADR-resistant breast cancer cells. *In vivo* experiments also confirmed that overexpression of GAS5 led to decreases in miR-221-3p and ABCB1 expression, as well as an increase in DKK2 expression, in resected tumors derived from MCF-7/ADR cells with or without ADR treatment. Taken together, all of these results led us to the conclusion that GAS5 could enhance ADR sensitivity in breast cancer cells via the miR-221-3p/DKK2 axis, at least in part through the Wnt/ β -catenin pathway.

In conclusion, our study demonstrates the existence of a dual role played by GAS5 in ABCB1-mediated drug resistance of breast cancer. In particular, we highlight a novel ceRNA-miRNA-mRNA regulation mechanism in which GAS5 positively reverses the ABCB1-mediated ADR resistance via the miR-221-3p/DKK2 axis by repressing the Wnt/ β -catenin pathway. Our present data provide strong evidence that the GAS5/miR-221-3p/DKK2 axis may be a promising chemosensitizing strategy for the treatment of breast cancer.

MATERIALS AND METHODS

Patients and Specimens

Twenty-six biopsy-proven patients with invasive primary breast cancer treated with NAC at the First Affiliated Hospital of the University of Science and Technology of China from January 2016 to December 2018 were enrolled in this study. The diagnosis of each case was confirmed by pathologists based on World Health Organization (WHO) classification.

Patients were treated with four cycles of epirubicin (90 or 100 mg/m²) combined with cyclophosphamide (600 mg/m²), followed by four cycles of docetaxel (90 or 100 mg/m²). Each chemotherapy cycle is 3 weeks. All patients received modified radical mastectomy after NAC. The data, including age, primary tumor size, lymph nodes metastasis, TNM (tumor, node, metastasis) stage, histological grade,

estrogen receptor (ER) status, progesterone receptor (PR) status, HER-2 status, and subtype, were collected from clinical and pathological records. The clinicopathological characteristics of the patients are summarized in Table 1. The clinical response was assessed by the decrease in tumor size and classified according to RECIST criteria.³³ Thus, patients with complete response (CR) or partial response (PR) were regarded as clinical responders, while patients with stable disease (SD) or progressive disease (PD) were regarded as non-responders for further analysis.^{34,35} Fresh tumor samples were immediately frozen in liquid nitrogen and preserved at -80°C . The present study was approved by the Ethics Committee of Anhui Medical University, and all patients signed an informed consent.

Cell Culture and Transfection

The parental-sensitive human breast cell line MCF-7 and ADR-resistant cell line MCF-7/ADR were kind gifts from Prof. Tao Zhu (University of Science and Technology of China). The cells were cultured in RPMI 1640 medium (Biological Industries, Israel) containing 10% fetal calf serum (Biological Industries, Israel) and 1% penicillin-streptomycin solution at 37°C under a humidified atmosphere of 5% CO_2 . The MCF-7/ADR cells were cultured in the above-mentioned media that were additionally supplemented with 1 $\mu\text{g}/\text{mL}$ ADR (KeyGen Biotech, Nanjing, China) to maintain the drug-resistant phenotype. The GAS5 overexpression plasmid pcDNA3.1-GAS5 (pcDNA-GAS5) was generated by inserting the full-length GAS5 sequence into the NheI and BamHI sites of pcDNA (Invitrogen, USA). All miRNA mimics (miR-221-3p NC [miR-NC]), miRNA inhibitors (anti-miR-221-3p, anti-miR-NC), and si-RNAs (si-NC, si-GAS5#1, and si-GAS5#2) were designed and synthesized by GenePharma (Generay, Shanghai, China). Transfection was performed using Lipofectamine 2000 (Invitrogen, USA) according to the manufacturer's instructions. Cells were harvested for further evaluation at 48 h after transfection. The sequences of oligonucleotides used are as follows: miR-221-3p mimics, 5'-AGCUACAUGUCUG CUGGGUUUC-3', 5'-AACCCAGCAGACAAUGUAGUCUUU-3'; miR-221-3p mimic NCs, 5'-UUCUCCGAACGUGUCACGUTT-3', 5'-ACGUGACACGUUCGGAGAATT-3'; miR-221-3p antagomirs, 5'-CAAACCCAGCAGACAAUGUAGCU-3'; miR-221-3p antagomir NCs, 5'-CAGUACUUUUGUAGUACAA-3'.

RNA-Seq Analysis

lncRNAs, miRNAs, and mRNAs in the RNA samples were profiled using human RNA-seq analysis (Smartquarier Biomedicine, Shanghai, China). Differentially expressed lncRNAs, miRNAs, and mRNAs were identified by volcano plot filtering. Upregulated or downregulated lncRNAs were selected based on changes ≥ 3 -fold thresholds and $p < 0.01$.

Drug Sensitivity Assay

The ADR sensitivity of cells was detected using a CCK-8 (Sigma-Aldrich) assay by measuring the IC_{50} value (ADR concentration causing 50% decrease in absorbance compared with control). Briefly, cells were seeded into each well of a 96-well plate at 2×10^3 per well overnight and incubated with various concentrations of ADR (0.02, 0.1,

0.5, 2.5, 12.5, and 25 $\mu\text{g}/\text{mL}$ for MCF-7, and 0.5, 1, 5, 25, 125, and 250 $\mu\text{g}/\text{mL}$ for MCF-7/ADR). After incubation with ADR treatment, the cell viability was evaluated using a CCK-8 assay according to the manufacturer's instructions.

Cell Proliferation Assay

Cell proliferation capacity was determined by using CCK-8 and colony formation assays. At 0, 24, 48, and 72 h after transfection, the CCK-8 solution was added to each well and incubated for 4 h. The absorbance was recorded at a wavelength of 450 nm using a SpectraMax i3x microplate reader (Molecular Devices, USA). With respect to the colony formation assay, cells were seeded in six-well plates and then cultured for 14 days in culture medium with 10% fetal bovine serum (FBS) as previously performed.³⁶ Each assay was repeated three times.

Apoptosis Analysis

Twenty-four hours after the transfection as described above, ADR was added, with the final concentration of 80 $\mu\text{g}/\text{mL}$ for MCF-7/ADR and 1 $\mu\text{g}/\text{mL}$ for MCF-7. Forty-eight hours later, cell apoptosis was measured by an annexin V-allophycocyanin (APC)/7-aminoactinomycin D (7-AAD) apoptosis detection kit (KeyGen Biotech, China) according to the manufacturer's protocols. Analyses were conducted using a BD LSR flow cytometer (BD Biosciences).

Intracellular Accumulation of Rh123

Accumulation of Rh123 by flow cytometry was carried out following modified methods as previously described.¹⁸ Briefly, the cells were transfected with GAS5 overexpression plasmid (pcDNA-GAS5) or GAS5 siRNA (si-GAS5#1) for 48 h, and then cells were collected and incubated with 5 $\mu\text{g}/\text{mL}$ Rh123 at 37°C , 5% CO_2 for 30 min. After incubation, the cells were gathered and washed three times with ice-cold PBS and then resuspended in 500 μL of PBS. Immediately, the mean fluorescence intensity of intracellular Rh123 was determined by a CytoFLEX flow cytometer (Beckman Coulter) with excitation/emission wavelengths of 488/525 nm.

Quantitative Real-Time PCR

Total RNA was extracted from breast cancer tissue samples or cell lines by using a RISO RNA isolation reagent (Biomics Biotechnologies, USA). The expression of miRNA was measured using an EzOmics miRNA qPCR detection primer set (Biomics Biotechnologies, USA) and EzOmics one-step qPCR kit (Biomics Biotechnologies, USA) in an Applied Biosystems 7500 instrument.

Total RNA was extracted by using TRIzol reagent (Invitrogen, USA). Reverse transcription was performed using a first-strand cDNA synthesis kit (Fermentas, USA) according to the manufacturer's protocol. Quantitative real-time PCR analysis was conducted by using SYBR Green PCR master mix (QIAGEN, Germany). The relative expression of target genes was calculated by using the $2^{-\Delta\Delta\text{CT}}$ method and expressed as fold change relative to the respective control. The CT values were normalized using U6 or GAPDH as an internal control. Primer sequences are listed in Table S1. PCR was performed in triplicate.

Separation of Nuclear and Cytoplasmic Fractions

The cytoplasmic and nuclear fractions were isolated from cells using a cytoplasmic and nuclear RNA purification kit (Norgen Biotek, Thorold, ON, Canada) as previously described.³⁷ Thus, the expression level of GAS5 in nuclear and cytoplasm fractions was detected by quantitative real-time PCR assays.

Western Blot Analysis

Cells were harvested and suspended in radioimmunoprecipitation assay (RIPA) lysis buffer (Beyotime, China) and quantified with a bicinchoninic acid (BCA) protein assay kit (Boster, Wuhan, China). After boiling in sample buffer, total proteins were separated by denaturing 8% or 10% sodium dodecyl sulfate-polyacrylamide gel electrophoresis (SDS-PAGE) and electrotransferred to polyvinylidene fluoride (PVDF) membranes (Millipore, Billerica, MA, USA). Subsequently, the membranes were then blocked and incubated with the following antibodies overnight at 4°C: anti-ABC1 (Millipore, Merck, Germany), anti- β -catenin, anti-c-Myc, and anti-cyclin D1 (Cell Signaling Technology, USA), anti-DKK2 (Abcam, England), and anti-GAPDH (Santa Cruz Biotechnology, Santa Cruz, CA, USA) as a control. Horseradish peroxidase-conjugated anti-mouse, anti-rabbit antibodies were used as secondary antibodies correspondingly. The bands were visualized by chemiluminescence reagents (ChemiQ 4600, Bioshine).

Luciferase Reporter Assay

Putative WT and MUT miR-221-3p-binding sites in the 3' UTR of GAS5 or DKK2 mRNA, termed GAS5-WT or GAS5-MUT and DKK2-WT or DKK2-MUT, were cloned into a pmirGLO-reporter luciferase vector (Promega, Fitchburg, WI, USA). The reporter plasmid was transiently transfected into MCF-7/ADR cells in the presence of miR-221-3p mimics, anti-miR-221-3p, and/or pcDNA-GAS5, si-GAS5#1. Forty-eight hours after transfection, relative luciferase activity was measured with the Dual-Luciferase reporter assay system (Promega, USA).

RIP Assay

RIP assays were performed using the EZ-Magna RIP kit (Millipore, Billerica, MA, USA) following the manufacturer's protocol. Briefly, the cells were lysed with RIP lysis buffer and then the cell extract was incubated with RIP buffer containing magnetic beads, which were conjugated with human anti-Ago2 antibody (Millipore) and NC normal mouse IgG (Millipore). Samples were then incubated with proteinase K buffer with shaking. Finally, the RNAs in the immunoprecipitates were extracted, purified, and detected by quantitative real-time PCR.

In Vivo Chemosensitivity Assay

The protocol was approved by the Animal Care and Ethics Committee of Anhui Medical University. All experimental procedures were performed in accordance with the National Institutes of Health animal use guidelines. Female BALB/c nude mice (4 weeks old, 18–20 g) were purchased from Slake Jingda Laboratory Animal Company (Hunan, China) and housed under specific pathogen-free

conditions. Approximately 2.0×10^7 MCF-7/ADR cells stably infected with pcDNA-GAS5 or pcDNA-control (NC) were subcutaneously injected into each mouse to form xenografts. When the tumor volume reached 50 mm³, ADR was intravenously injected via the tail at a dose of 5 mg/kg every 4 days for five times according to the indicated groups (n = 6 for each group): pcDNA-NC+PBS, pcDNA-NC+ADR, pcDNA-GAS5+PBS, and pcDNA-GAS5+ADR. Tumor growth was examined every 4 days, and tumor volumes were calculated with the following equation: volume = $0.5 \times \text{width}^2 \times \text{length}$. After 27 days, the mice were all sacrificed and the tumor masses were dissected, photographed, and subjected to statistical analyses.

Statistical Analysis

Data were expressed as the mean \pm SD and analyzed using SPSS 19.0 statistics software (SPSS, USA). Significant differences among groups were determined by a Student's t test or one-way ANOVA. Correlations between GAS5 and miR-221-3p were analyzed by Spearman rank correlation. The statistical significance was defined as *p < 0.05 or **p < 0.01. All experiments were performed at least three times.

SUPPLEMENTAL INFORMATION

Supplemental Information can be found online at <https://doi.org/10.1016/j.omtn.2020.01.030>.

AUTHOR CONTRIBUTIONS

Study Design, A.S. and L.Z.; Drafting the Manuscript, Z.C.; Critical Revision of the Manuscript, D.J. and L.J.; Performance of Experiments, Z.C., T.P., and D.J.; Preparation and Assistant Work, Y.G. and X.F.; Necessary Technical Guides, L.Z. and X.F.

CONFLICTS OF INTEREST

The authors declare no competing interests.

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REFERENCES

1. Siegel, R.L., Miller, K.D., and Jemal, A. (2019). Cancer statistics, 2019. *CA Cancer J. Clin.* 69, 7–34.
2. Li, Y., Gao, X., Yu, Z., Liu, B., Pan, W., Li, N., and Tang, B. (2018). Reversing multi-drug resistance by multiplexed gene silencing for enhanced breast cancer chemotherapy. *ACS Appl. Mater. Interfaces* 10, 15461–15466.
3. Chang, L., Hu, Z., Zhou, Z., and Zhang, H. (2018). Linc00518 contributes to multi-drug resistance through regulating the mir-199a/MRP1 axis in breast cancer. *Cell. Physiol. Biochem.* 48, 16–28.
4. Yao, N., Fu, Y., Chen, L., Liu, Z., He, J., Zhu, Y., Xia, T., and Wang, S. (2019). Long non-coding RNA NONHSAT101069 promotes epirubicin resistance, migration, and

- invasion of breast cancer cells through NONHSAT101069/miR-129-5p/Twist1 axis. *Oncogene* 38, 7216–7233.
5. Chen, Z., Ma, T., Huang, C., Zhang, L., Lv, X., Xu, T., Hu, T., and Li, J. (2013). miR-27a modulates the MDR1/P-glycoprotein expression by inhibiting FZD7/ β -catenin pathway in hepatocellular carcinoma cells. *Cell. Signal.* 25, 2693–2701.
 6. Tian, F., Dahmani, F.Z., Qiao, J., Ni, J., Xiong, H., Liu, T., Zhou, J., and Yao, J. (2018). A targeted nanoplatform co-delivering chemotherapeutic and antiangiogenic drugs as a tool to reverse multidrug resistance in breast cancer. *Acta Biomater.* 75, 398–412.
 7. Hu, J., Wang, Z., Shan, Y., Pan, Y., Ma, J., and Jia, L. (2018). Long non-coding RNA HOTAIR promotes osteoarthritis progression via miR-17-5p/FUT2/ β -catenin axis. *Cell Death Dis.* 9, 711.
 8. Dong, H., Hu, J., Zou, K., Ye, M., Chen, Y., Wu, C., Chen, X., and Han, M. (2019). Activation of lncRNA TINCR by H3K27 acetylation promotes trastuzumab resistance and epithelial-mesenchymal transition by targeting microRNA-125b in breast cancer. *Mol. Cancer* 18, 3.
 9. Han, J., Han, B., Wu, X., Hao, J., Dong, X., Shen, Q., and Pang, H. (2018). Knockdown of lncRNA H19 restores chemo-sensitivity in paclitaxel-resistant triple-negative breast cancer through triggering apoptosis and regulating Akt signaling pathway. *Toxicol. Appl. Pharmacol.* 359, 55–61.
 10. Gao, H., Hao, G., Sun, Y., Li, L., and Wang, Y. (2018). Long noncoding RNA H19 mediated the chemosensitivity of breast cancer cells via Wnt pathway and EMT process. *OncoTargets Ther.* 11, 8001–8012.
 11. Dong, H., Wang, W., Mo, S., Liu, Q., Chen, X., Chen, R., Zhang, Y., Zou, K., Ye, M., He, X., et al. (2018). Long non-coding RNA SNHG14 induces trastuzumab resistance of breast cancer via regulating *PABPC1* expression through H3K27 acetylation. *J. Cell. Mol. Med.* 22, 4935–4947.
 12. Dong, H., Wang, W., Chen, R., Zhang, Y., Zou, K., Ye, M., He, X., Zhang, F., and Han, J. (2018). Exosome-mediated transfer of lncRNA-SNHG14 promotes trastuzumab chemoresistance in breast cancer. *Int. J. Oncol.* 53, 1013–1026.
 13. Schneider, C., King, R.M., and Philipson, L. (1988). Genes specifically expressed at growth arrest of mammalian cells. *Cell* 54, 787–793.
 14. Arshi, A., Sharifi, F.S., Khorramian Ghahfarokhi, M., Faghih, Z., Doosti, A., Ostovari, S., Mahmoudi Maymand, E., and Ghahramani Seno, M.M. (2018). Expression analysis of *MALAT1*, *GAS5*, *SRA*, and *NEAT1* lncRNAs in breast cancer tissues from young women and women over 45 years of age. *Mol. Ther. Nucleic Acids* 12, 751–757.
 15. Mourtada-Maarabouni, M., Pickard, M.R., Hedge, V.L., Farzaneh, F., and Williams, G.T. (2009). *GAS5*, a non-protein-coding RNA, controls apoptosis and is downregulated in breast cancer. *Oncogene* 28, 195–208.
 16. Gu, J., Wang, Y., Wang, X., Zhou, D., Shao, C., Zhou, M., and He, Z. (2018). Downregulation of lncRNA *GAS5* confers tamoxifen resistance by activating miR-222 in breast cancer. *Cancer Lett.* 434, 1–10.
 17. Zhang, Z., Zhu, Z., Watabe, K., Zhang, X., Bai, C., Xu, M., Wu, F., and Mo, Y.Y. (2013). Negative regulation of lncRNA *GAS5* by miR-21. *Cell Death Differ.* 20, 1558–1568.
 18. Chen, Z., Huang, C., Ma, T., Jiang, L., Tang, L., Shi, T., Zhang, S., Zhang, L., Zhu, P., Li, J., et al. (2018). Reversal effect of quercetin on multidrug resistance via FZD7/ β -catenin pathway in hepatocellular carcinoma cells. *Phytomedicine* 43, 37–45.
 19. Ye, K., Wang, S., Zhang, H., Han, H., Ma, B., and Nan, W. (2017). Long noncoding RNA *GAS5* suppresses cell growth and epithelial-mesenchymal transition in osteosarcoma by regulating the miR-221/ARHI pathway. *J. Cell. Biochem.* 118, 4772–4781.
 20. Xiang, S., Dauchy, R.T., Hoffman, A.E., Pointer, D., Frasca, T., Blask, D.E., and Hill, S.M. (2019). Epigenetic inhibition of the tumor suppressor ARHI by light at night-induced circadian melatonin disruption mediates STAT3-driven paclitaxel resistance in breast cancer. *J. Pineal Res.* 67, e12586.
 21. Perone, Y., Farrugia, A.J., Rodríguez-Meira, A., Györfy, B., Ion, C., Uggetti, A., Chronopoulos, A., Marrazzo, P., Faronato, M., Shousha, S., et al. (2019). SREBP1 drives Keratin-80-dependent cytoskeletal changes and invasive behavior in endocrine-resistant ER α breast cancer. *Nat. Commun.* 10, 2115.
 22. Díaz-Rodríguez, E., Pérez-Peña, J., Ríos-Luci, C., Arribas, J., Ocaña, A., and Pandiella, A. (2019). TRAIL receptor activation overcomes resistance to trastuzumab in HER2 positive breast cancer cells. *Cancer Lett.* 453, 34–44.
 23. Zhang, J., Du, Z., Pan, S., Shi, M., Li, J., Yang, C., Hu, H., Qiao, M., Chen, D., and Zhao, X. (2018). Overcoming multidrug resistance by codelivery of MDR1-targeting siRNA and doxorubicin using EphA10-mediated pH-sensitive lipoplexes: in vitro and in vivo evaluation. *ACS Appl. Mater. Interfaces* 10, 21590–21600.
 24. Liu, B., Wu, S., Ma, J., Yan, S., Xiao, Z., Wan, L., Zhang, F., Shang, M., and Mao, A. (2018). lncRNA *GAS5* reverses EMT and tumor stem cell-mediated gemcitabine resistance and metastasis by targeting miR-221/SOCS3 in pancreatic cancer. *Mol. Ther. Nucleic Acids* 13, 472–482.
 25. Yao, T., Lu, R., Zhang, J., Fang, X., Fan, L., Huang, C., Lin, R., and Lin, Z. (2019). Growth arrest-specific 5 attenuates cisplatin-induced apoptosis in cervical cancer by regulating STAT3 signaling via miR-21. *J. Cell. Physiol.* 234, 9605–9615.
 26. Chen, Z., Shi, T., Zhang, L., Zhu, P., Deng, M., Huang, C., Hu, T., Jiang, L., and Li, J. (2016). Mammalian drug efflux transporters of the ATP binding cassette (ABC) family in multidrug resistance: a review of the past decade. *Cancer Lett.* 370, 153–164.
 27. Deng, L., Lei, Q., Wang, Y., Wang, Z., Xie, G., Zhong, X., Wang, Y., Chen, N., Qiu, Y., Pu, T., et al. (2017). Downregulation of miR-221-3p and upregulation of its target gene *PARP1* are prognostic biomarkers for triple negative breast cancer patients and associated with poor prognosis. *Oncotarget* 8, 108712–108725.
 28. Ergun, S., Tayeb, T.S., Arslan, A., Temiz, E., Arman, K., Safdar, M., Dağlı, H., Korkmaz, M., Nacarkahya, G., Kırkeş, S., and Oztuzcu, S. (2015). The investigation of miR-221-3p and *PAK1* gene expressions in breast cancer cell lines. *Gene* 555, 377–381.
 29. Li, G.Y., Wang, W., Sun, J.Y., Xin, B., Zhang, X., Wang, T., Zhang, Q.F., Yao, L.B., Han, H., Fan, D.M., et al. (2018). Long non-coding RNAs AC026904.1 and UCA1: a “one-two punch” for TGF- β -induced SNAI2 activation and epithelial-mesenchymal transition in breast cancer. *Theranostics* 8, 2846–2861.
 30. Deng, F., Zhou, R., Lin, C., Yang, S., Wang, H., Li, W., Zheng, K., Lin, W., Li, X., Yao, X., et al. (2019). Tumor-secreted dickkopf2 accelerates aerobic glycolysis and promotes angiogenesis in colorectal cancer. *Theranostics* 9, 1001–1014.
 31. Wang, Y., Zhao, Y., Herbst, A., Kalinski, T., Qin, J., Wang, X., Jiang, Z., Benedix, F., Franke, S., Wartman, T., et al. (2016). miR-221 mediates chemoresistance of esophageal adenocarcinoma by direct targeting of *DKK2* expression. *Ann. Surg.* 264, 804–814.
 32. Xiao, Q., Wu, J., Wang, W.J., Chen, S., Zheng, Y., Yu, X., Meeth, K., Sahraei, M., Bothwell, A.L.M., Chen, L., et al. (2018). *DKK2* imparts tumor immunity evasion through β -catenin-independent suppression of cytotoxic immune-cell activation. *Nat. Med.* 24, 262–270.
 33. Therasse, P., Arbuck, S.G., Eisenhauer, E.A., Wanders, J., Kaplan, R.S., Rubinstein, L., Verweij, J., Van Glabbeke, M., van Oosterom, A.T., Christian, M.C., and Gwyther, S.G. (2000). New guidelines to evaluate the response to treatment in solid tumors. European Organization for Research and Treatment of Cancer, National Cancer Institute of the United States, National Cancer Institute of Canada. *J. Natl. Cancer Inst.* 92, 205–216.
 34. Paik, J.H., Jang, J.Y., Jeon, Y.K., Kim, W.Y., Kim, T.M., Heo, D.S., and Kim, C.W. (2011). MicroRNA-146a downregulates NF κ B activity via targeting TRAF6 and functions as a tumor suppressor having strong prognostic implications in NK/T cell lymphoma. *Clin. Cancer Res.* 17, 4761–4771.
 35. Chintamani, S., Singh, J.P., Mittal, M.K., Saxena, S., Bansal, A., Bhatia, A., and Kulshreshtha, P. (2005). Role of p-glycoprotein expression in predicting response to neoadjuvant chemotherapy in breast cancer—a prospective clinical study. *World J. Surg. Oncol.* 3, 61.
 36. Song, X., Zhang, X., Wang, X., Chen, L., Jiang, L., Zheng, A., Zhang, M., Zhao, L., and Wei, M. (2020). lncRNA *SPRY4-IT1* regulates breast cancer cell stemness through competitively binding miR-6882-3p with TCF7L2. *J. Cell. Mol. Med.* 24, 772–784.
 37. Zaghlool, A., Ameer, A., Nyberg, L., Halvardson, J., Grabherr, M., Cavelier, L., and Feuk, L. (2013). Efficient cellular fractionation improves RNA sequencing analysis of mature and nascent transcripts from human tissues. *BMC Biotechnol.* 13, 99.