Aberrant epigenetic regulation of GABRP associates with aggressive phenotype of ovarian cancer

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Metastasis is a major cause of therapeutic failure in ovarian cancer. To elucidate molecular mechanisms of ovarian cancer metastasis, we previously established a metastatic xenograft mouse model using human ovarian carcinoma SK-OV-3 cells. Using gene expression profiling, we found that γ-aminobutyric acid (GABA)A receptor π subunit (GABRP) expression was upregulated (> 4-fold) in metastatic tissues from our xenograft mice compared with SK-OV-3 cells. Importantly, GABRP knockdown diminished the migration and invasion of SK-OV-3 cells, and reduced extracellular signal-regulated kinase (ERK) activation while overexpression of GABRP exhibited significantly increased cell migration, invasion and ERK activation. Moreover, treatment with the mitogen-activated protein kinase (MAPK)/ERK kinase (MEK) inhibitor U0126 similarly suppressed the migration and invasion of SK-OV-3 cells, implying that GABRP promotes these cellular behaviors by activating the MAPK/ERK pathway. Using genome-wide DNA methylation profiling, we identified hypomethylated CpG sites in the GABRP promoter in metastatic tissues from the xenograft mice compared with SK-OV-3 cells. Treatment with a DNA methyltransferase inhibitor demonstrated that methylation at −963 bp from the GABRP transcription start site (−963 CpG site) was critical for the epigenetic regulation of GABRP. Finally, we analyzed human ovarian cancer patient samples and showed DNA hypomethylation at the GABRP −963 CpG site in advanced stage, but not early-stage, primary tumors compared with their paired normal tissues. These findings suggest that GABRP enhances the aggressive phenotype of ovarian cancer cells, and that the DNA methylation status of the GABRP −963 CpG site may be useful for predicting the metastatic potential in ovarian cancer patients.

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INTRODUCTION

Ovarian cancer has the highest mortality rate among gynecological cancers, with a 5-year survival rate of only 27% when the disease is diagnosed at an advanced stage.1 The majority of ovarian cancer patients are diagnosed at an advanced stage because they show few clear symptoms, and there are no reliable diagnostic tools for early detection. Ovarian cancer is unique in that it mainly metastasizes via a transcoelomic route leading to ascites-mediated intraperitoneal dissemination rather than hematogenous dissemination.2 The intraperitoneal dissemination occurs relatively early in the disease, which is another reason why ovarian cancer is rarely diagnosed at an early stage.3 Because metastasis prevention is the major obstacle in the clinical management of ovarian cancer, a comprehensive understanding of the molecular mechanisms that regulate the aggressive behavior of ovarian cancer cells is required to improve treatment outcomes for ovarian cancer patients.

We previously injected human ovarian carcinoma SK-OV-3 cells into the peritoneum of female nude mice to establish an ovarian cancer xenograft mouse model.4 We then performed expression profiling to identify genes that were differentially expressed in the metastatic tissues, compared with the injected SK-OV-3 cells.4–7 Among the genes whose expression was altered more than twofold in metastatic tissues, we selected γ-aminobutyric acid (GABA)A receptor π subunit (GABRP), which was upregulated in the metastatic tissue, for further study.

GABA is the major inhibitory neurotransmitter in the mature mammalian central nervous system; it functions through activation of the GABA receptors, which include the ionotropic (GABA_A or GABA_C) and the metabotropic
(GABA<sub>B</sub>)<sup>8,9</sup> receptors. The GABA<sub>A</sub> receptor is a heteropentameric, ligand-gated chloride channel comprised of various subunits (α1–6, β1–3, γ1–3, δ, ε, θ, π and ρ1–3).<sup>10,11</sup>

GABA and GABA receptors are also expressed in peripheral non-neuronal cells, suggesting that they have additional functions beyond regulating neurotransmission. GABA and GABA receptors have been reported to regulate proliferation, differentiation, migration and development of many peripheral non-neuronal cells; however, their precise roles in non-neuronal cells, including cancer cells, are largely unknown.<sup>12</sup> Conversely, GABRP is mainly expressed in non-neuronal tissues, including the mammary gland, prostate gland, lung, thymus and uterus; it is not found abundantly in the brain.<sup>13,14</sup> GABRP is also highly expressed in certain types of cancer such as pancreatic ductal adenocarcinoma<sup>15</sup> and basal-like breast cancer.<sup>16</sup> In pancreatic ductal adenocarcinoma, GABRP promotes proliferation by increasing intracellular Ca<sup>2+</sup> levels and activating the mitogen-activated protein kinase/extracellular signal-regulated kinase (MAPK/ERK) signaling pathway.<sup>15</sup> In basal-like breast cancer cells, GABRP stimulates the migratory activity and expression of basal-like cytokeratins through ERK cascade.<sup>16</sup>

On the basis of these previously established roles of GABRP in tumorigenesis and our identification of GABRP as an upregulated gene in metastatic tissues compared with ovarian carcinoma cells from our xenograft mouse model, we hypothesized that high GABRP expression in ovarian carcinoma cells actively enhances their metastatic potential. Thus, in this study, we performed GABRP gain- and loss-of-function studies in the SK-OV-3 ovarian carcinoma cell line to investigate its role in ovarian cancer metastasis. Moreover, we analyzed the DNA methylation status of the GABRP promoter using our previously established ovarian cancer xenograft mouse model to determine whether the differential expression of GABRP in metastatic tissues is epigenetically regulated. Finally, we analyzed ovarian cancer patient samples to determine whether GABRP expression and/or epigenetic modifications are clinically relevant.

**MATERIALS AND METHODS**

**Cell culture**

The human ovarian cancer cell line SK-OV-3 was purchased from American Type Culture Collection (ATCC, no. HTB-77, Manassas, VA, USA) and cultured in McCoy’s 5A medium (Gibco/BRL, Rockville, MD, USA) supplemented with 10% fetal bovine serum (FBS, Gibco/BRL), 100 U ml<sup>−1</sup> penicillin (Gibco/BRL) and 100 μg ml<sup>−1</sup> streptomycin (Gibco/BRL) in an atmosphere of 95% humidified air and 5% CO<sub>2</sub> at 37°C.

**Ovarian cancer xenograft mouse model**

All procedures for handling and killing the animals used in this study were performed in strict compliance with the guidelines of the Korean animal protection law, and were approved by the Institutional Animal Care and Use Committee (IACUC) of Ewha Womans University School of Medicine. SK-OV-3 cells (2 × 10<sup>6</sup>) suspended in culture medium were injected intraperitoneally into 10 female nude mice (CAnN.Cg-Foxn1<sup>nu</sup>, 4–6 weeks old). Four weeks after inoculation, the xenograft mice were killed, and at least four tumor metastases adhering to the mesothelial surface of the peritoneum were harvested from each mouse.

**Processing of mRNA microarray and gene expression analysis**

Total RNA was extracted from the tumor metastases of the mice and SK-OV-3 cells using the RNeasy Mini Kit (Qiagen, Valencia, CA, USA). One microgram of total RNA was amplified and labeled according to the Affymetrix GeneChip Whole Transcript Sense Target Labeling protocol. The labeled cDNA was hybridized to Affymetrix Human Gene 1.0 ST arrays (Affymetrix, Santa Clara, CA, USA). The scanned raw expression values were background-corrected, normalized and summarized using the Robust Multiarray Averaging approach in the Bioconductor ‘affy’ package (Affymetrix). The resulting log<sub>2</sub>-transformed data were used for further analyses.

To identify differentially expressed genes (DEGs), we applied moderated t-statistics based on an empirical Bayesian approach.<sup>17</sup> Significantly upregulated and downregulated DEGs were defined as genes with at least a twofold difference in expression levels between the xenograft tissues and SK-OV-3 cells after multiple testing correction (Benjamini–Hochberg false-discovery rate-adjusted P-value < 0.05).<sup>18</sup> Finally, we excluded genes with a low expression level (maximum log<sub>2</sub> expression level in a total of eight samples < 7.0) from the list of DEGs.

**RNA preparation and reverse-transcription quantitative polymerase chain reaction**

Total RNA was extracted from the tumor metastases and SK-OV-3 cells using the RNeasy Mini Kit (Qiagen) according to the manufacturer’s protocol. One microgram of total RNA was converted to cDNA using Superscript II reverse transcriptase (Invitrogen, Carlsbad, CA, USA) and oligo-(dT)<sub>12-18</sub> primers (Invitrogen) according to the manufacturer’s instructions. Reverse-transcription quantitative polymerase chain reaction (RT-qPCR) was performed in a 20 μl reaction mixture containing 1 μl cDNA, 10 μl SYBR Premix EX Taq (Takara Bio, Otsu, Japan), 0.4 μl Rox reference dye (50 x, Takara Bio), and 200 nm primers for each gene. The primer sequences were: GABRP (forward), 5′-CTCGATTTCTCCTGGCA-3′; GABRP (reverse), 5′-GTGCGGTCACCGGATC-3′; GAPDH (forward), 5′-AATGCTTCACACATGAT-3′; and GAPDH (reverse), 5′-TGAGTCTGAGTGTTCTG-3′. The reactions were run on a 7500 Fast Real-Time PCR System (Applied Biosystems, Foster City, CA, USA) at 95 °C for 30 s, followed by 40 cycles of 95 °C for 3 s and 60 °C for 30 s and a single cycle of 95 °C for 15 s, 60 °C for 30 s, and 95 °C for 15 s to generate dissociation curves. All PCR reactions were performed in triplicate, and the specificity of the reaction was determined by melting curve analysis. Comparative quantification of each target gene was performed based on cycle threshold (C<sub>T</sub>) normalized to GAPDH using the ΔΔC<sub>T</sub> method.

**Genomic DNA isolation and CpG methylation microarray**

Genomic DNA was extracted from cell lines and tumor metastases using a QIAamp mini kit (Qiagen) according to the manufacturer’s instructions. The Illumina HumanMethylation450 BeadChip (Illumina, San Diego, CA, USA) that targets ~ 450 000 CpG sites was used for genome-wide screening of DNA methylation. DNA methylation values were described as β-values, which were calculated by subtracting the background using negative controls on the array and taking the ratio of the methylated signal intensity against the sum
of both methylated and unmethylated signals. The $\beta$-values ranged from 0 (completely unmethylated) to 1 (fully methylated) on a continuous scale for each CpG site. To identify differentially methylated CpG sites, we applied the difference in mean $\beta$-values ($\Delta\beta$; mean $\beta$-value in tumors minus mean $\beta$-value in SK-OV-3). If the absolute difference of the mean $\beta$-values ($|\Delta\beta|$) was >0.2, the site was defined as a differentially methylated CpG site. We described a CpG site as hypermethylated if $\Delta\beta$ was greater than 0.2 and hypomethylated if $\Delta\beta$ was less than −0.2.

**Bisulfite sequencing PCR**

Genomic DNA was extracted from the harvested tumor metastases of the ovarian cancer mouse xenografts and SK-OV-3 cells using the QIAamp DNA mini kit (Qiagen) according to the manufacturer's protocol. Bisulfite treatment of the genomic DNA was performed using the EpiTect Bisulfite Kit (Qiagen) according to the manufacturer's instructions. For bisulfite sequencing of the target promoter region of **GABRP**, bisulfite sequencing PCR (BSP) was carried out using conventional PCR in a 50 µl reaction mixture containing 10 ng bisulfite-modified genomic DNA, 1.5 mM MgCl$_2$, 200 µM dNTP, 1 U Platinum Taq polymerase (Invitrogen), 1 × Platinum Taq buffer and 200 nM gene-specific BSP forward and reverse primers. The BSP primers were designed using MethPrimer software (http://www.urogene.org/methprimer). For **GABRP**, the BSP product was 452 bp (position in the human GRC37/hg19 assembly: chromosome 5, 170209520–170209971) and contained five CpG sites. The five CpG sites are located at −1178, −1142, −994, −963 and −924 from the transcription start site. The BSP primer sequences were: (forward), 5′-TGAATCTTTTATTAGGAGGAAATGAG-3′; and (reverse), 5′-CTAAACTCCTATCCCTACCT-3′. The reaction ran at 95 °C for 5 min, followed by 30 cycles of 95 °C for 30 s, 55 °C for 30 s and 72 °C for 30 s, and a final elongation step at 72 °C for 5 min.

The BSP products were purified using the QIAquick Gel Extraction kit (Qiagen) according to the manufacturer's protocols and ligated into the yT&A cloning vector (Yeastern Biotech, Taipei, Taiwan). The ligation products were used to transform competent DH5α Escherichia coli cells (RBC Bioscience, Taipei, Taiwan) using standard procedures. Blue/white screening was used to select bacterial clones, and BSP product-positive clones were confirmed by colony PCR using the BSP primers to verify the insert plasmid. Genomic DNA was then extracted from at least 15 insert-positive clones using the QIAprep Spin Miniprep kit (Qiagen) and sequenced using the M13 primer to analyze the methylation status at specific CpG sites.

**Quantitative methylation-specific PCR (qMSP)** was carried out with bisulfite-modified genomic DNA as the template and specific primer sequences designed to detect the methylated and unmethylated forms of two CpGs at −1142 and +963 bp from the transcription start site. The following methylated/unmethylated-specific primers were used: −1142 CpG (methylated forward), 5′-GGTCTTTATTAGGAGGAAATGAG-3′; −1142 CpG (unmethylated forward), 5′-GGTCTTTATTAGGAGGAAATGAG-3′; −1142 CpG (reverse), 5′-AAACTCCTATCCCTACCT-3′; −963 CpG (methylated forward), 5′-GATTTCTTTATTAGGAGGAAATGAG-3′; −963 CpG (reverse), 5′-AAACTCTATTAGGAGGAAATGAG-3′. For qMSP, a 20 µl reaction mixture containing 2 µl (10–100 ng µl$^{-1}$) bisulfite-treated DNA, 10 µl SYBR Premix EX Taq (Takara Bio), 0.4 µl Rox reference dye (50×; Takara Bio), and 200 nM each primer was amplified using a 7500 Fast Real-Time PCR system (Applied Biosystems). The amplification reaction conditions were 95 °C for 30 s, followed by 40 cycles of 95 °C for 3 s and 63 °C for 30 s. The PCR product was then cycled at 95 °C for 15 s, 60 °C for 1 min, and 95 °C for 15 s to generate the dissociation curves for analysis. The methylation percentage at each CpG site was calculated as follows ($C$ represents the threshold cycle):

$$\%\text{Methylation} = \frac{100}{1 + 2^{\left(\frac{C_{\text{unmeth}} - C_{\text{meth}}}{\Delta C_{\text{t}}}ight)}}$$

**5-aza-2′-deoxycytidine treatment**

To demethylate methylated CpG sites, SK-OV-3 cells were treated with an increasing concentration (0, 5, and 10 μM) of 5-aza-2′-deoxycytidine (5-aza-dC; Sigma-Aldrich, St Louis, MO, USA) for 3 days. The culture medium was replaced daily.

**Transient transfection**

To establish a transient expression system, SK-OV-3 cells were transfected with pCMV6-GABRP (Origene, Rockville, MD, USA) or pEGFP-N3 (Clontech, Mountain View, CA, USA) plasmids using Lipofectamine 2000 (Invitrogen). Briefly, cells were plated at 6 × 10$^5$ cells per well in six-well plates and allowed to grow overnight. Two micrograms of each plasmid and 5 μl of Lipofectamine 2000 were diluted in Opti-MEM medium (Gibco/BRL) to a total volume of 250 μl. The plasmid-Lipofectamine 2000 mixture was incubated at room temperature for 20 min. The transfection mixture was added to the six-well plates containing complete growth medium, and plates were incubated at 37 °C for 24 h in a 5% CO$_2$ incubator. Overexpression of **GABRP** was confirmed by qPCR 24 h post transfection.

Pre-designed small interfering RNA (siRNA) for **GABRP** (siGABRP, CAT# ID LU-006176-00-0005) and a non-targeting control (siNC, CAT# ID D-001206-13-05) were purchased from GE Healthcare (Lafayette, CO, USA). To deplete **GABRP**, SK-OV-3 cells were transfected with 100 nM siGABRP or siNC using DharmaFECT 1 transfection reagent (GE Healthcare) according to the manufacturer’s protocol. Knockdown of **GABRP** was confirmed using RT-qPCR 24 h post transfection.

**Transwell migration and in vitro invasion assay**

Complete growth medium was changed to serum-free medium 24 h post transfection. After 24 h of serum deprivation, cell migration assays were performed in 24-well transwell plates with 8-µm pore size membrane inserts (Corning, New York, NY, USA) as previously described.° SK-OV-3 cells in serum-free medium, with or without the MAPK/ERK kinase (MEK) inhibitor U0126 (10 µM) (Cell Signaling Technology, Danvers, MA, USA), were added at 2 × 10$^5$ cells per well into the upper chambers of the inserts, whereas medium containing 15% FBS was added to the lower chambers as a chemoattractant. For each experiment, both chemotactic migration to medium containing 15% FBS and random migration in serum-free medium were assessed in parallel transwell plates for 6 h at 37 °C in a 5% CO$_2$ incubator. The **in vitro** invasion assays were performed using the BD BioCoat Matrigel Invasion Chamber kit (Becton-Dickinson, Franklin Lakes, NJ, USA) as previously described.° SK-OV-3 cells in serum-free medium, with or without 10 µM U0126 (Cell Signaling Technology), were seeded at 1 × 10$^5$ cells per well into the upper chambers, whereas the medium containing 10% FBS was added to the lower chambers. The migration through the Matrigel chamber was allowed to proceed at 37 °C for 24 h in a 5% CO$_2$ incubator.
After the incubation periods for cellular migration or invasion, the SK-OV-3 cells on the upper side of the filter that had not migrated or invaded were carefully scraped away with cotton swabs, whereas those on the lower side were fixed for 2 min using the Diff-Quick kit solution (Thermo Fisher Scientific, Waltham, MA, USA), stained with 1% crystal violet for 2 min, and washed twice with distilled water at room temperature. The images of stained cells were acquired at ×200 magnification from six different fields. For quantitative analysis, the stained cells were subsequently extracted with 10% acetic acid, and colorimetric measurements were performed at 590 nm.

**Western blot analyses**

SK-OV-3 cell lines were lysed in RIPA buffer (Sigma-Aldrich) containing proteinase inhibitor cocktail (Roche Applied Science, Mannheim, Germany). Proteins (20 μg) were resolved using denaturing 10% sodium dodecyl sulfate-polyacrylamide gel electrophoresis and transferred to polyvinylidene fluoride membranes. Membranes were blocked in 5% skim milk in Tris-buffered saline with 0.1% Tween 20 (TBST) and subsequently incubated overnight at 4 °C with the following primary antibodies: p44/42 MAPK (Erk1/2) (137F5) rabbit mAb, and subsequently incubated overnight at 4 °C with the following secondary antibodies conjugated to horseradish peroxidase for 1 h at room temperature. After washing, the membranes were incubated with secondary antibodies conjugated to horseradish peroxidase for 1 h at room temperature. Protein bands were detected via chemiluminescence (ECL) using a ChemiDoc imaging system (Bio-Rad, Hercules, CA, USA). Western blot analyses were performed under normal conditions.

**Tissue specimen collection**

Six patients with early stage (stages I and II), and nine patients with advanced stage, ovarian cancer were included in this study. Primary solid tumor and adjacent normal tissues were collected from the Ewha Biospecimen Bank. Patient information, including age, sex, bHistological grade.

| Patients no. | Age | Stage | Grade | Histology                      |
|--------------|-----|-------|-------|--------------------------------|
| #1           | 57  | IIC   | G3    | Endometrioid adenocarcinoma    |
| #2           | 53  | IVA   | G2    | Serous adenocarcinoma          |
| #3           | 64  | IIC   | G3    | Serous papillary carcinoma     |
| #4           | 70  | IIC   | G3    | Serous papillary carcinoma     |
| #5           | 48  | IIC   | G3    | Serous carcinoma               |
| #6           | 51  | IV    | G3    | Serous papillary carcinoma     |
| #7           | 42  | I VB  | G3    | Serous papillary carcinoma     |
| #8           | 54  | IIC   | G3    | Serous carcinoma               |
| #9           | 55  | IIC   | G3    | Serous carcinoma               |
| #10          | 45  | IC1   | G3    | Clear cell carcinoma           |
| #11          | 49  | Ia    | G2    | Endometrioid adenocarcinoma    |
| #12          | 54  | IC2   | GX    | Mucinous borderline tumor      |
| #13          | 42  | IIB   | G3    | Mixed endometrioid adenocarcinoma and serous adenocarcinoma |
| #14          | 38  | Ia    | GX    | Clear cell carcinoma           |
| #15          | 52  | Ia    | G2    | Clear cell carcinoma           |

*International Federation of Gynecology and Obstetrics (FIGO) stage.*

*Histological grade.*
A recent study reported that GABRP was required for maintaining the pro-migratory behavior of basal-like breast cancer cells via activation of the ERK signaling pathway.\(^{16}\) Therefore, we investigated whether the ERK signaling pathway is involved downstream of GABRP in promoting the aggressive behavior of ovarian cancer cells. Indeed, SK-OV-3 cells treated with the MAPK/ERK kinase (MEK) inhibitor U0126 showed a 64% reduction in migration, and a 39% reduction in invasion, compared with those of untreated cells (Figure 2c).

Furthermore, immunoblot analyses confirmed that ERK1/2 activation was effectively inhibited by U0126 and showed that overexpression of GABRP enhanced ERK1/2 activation while GABRP knockdown diminished ERK1/2 activation (Figure 2d) indicating that GABRP promotes the metastatic potential of ovarian carcinoma cells through ERK pathway activation. Next, we examined whether alteration of GABRP expression could affect intracellular calcium signaling. SK-OV-3 cells were transfected with EGFP and GABRP expression constructs or with siNC and siGABRP. After 24 h transfection, the cells were treated with 100 μM muscimol, a GABA\(_A\) receptor agonist. Intracellular calcium concentration was determined using a colorimetric calcium detection kit. Although muscimol treatment increased intracellular calcium concentrations by 1.96-fold, overexpression of GABRP did not induce intracellular calcium mobilization (Supplementary Table S1). These data suggest that it was unlikely that calcium influx through voltage-dependent calcium channel affects the GABRP-induced metastatic potential in ovarian carcinoma cells.

The GABRP promoter region was hypomethylated in metastatic tissues from ovarian cancer xenograft mice

Previously, we performed genome-wide DNA methylation profiling using CpG methylation microarrays to examine the alterations in DNA methylation between metastatic tissues from ovarian cancer xenograft mice and ovarian carcinoma cells.\(^5\)–\(^7\) Using these data, we identified two CpG sites within the GABRP promoter region that were clearly hypomethylated in all tested metastatic tissues compared with those of the ovarian carcinoma cells. DNA methylation at these two CpG sites, located at −1142 and −963 bp from the transcription start site, was decreased by 20% and 59%, respectively, in the metastatic tissues compared with those of the ovarian carcinoma cells (Figures 3a and b). Next, we performed qMSP and confirmed that DNA methylation at −1142 and −963 CpG sites was reduced approximately 5–13% and 36–56%, respectively, in the metastatic tissues compared with those of the SK-OV-3 cells (Figures 3c and d).

To further analyze the DNA methylation status of the GABRP promoter region, which is located at 170209520–170209971 on chromosome 5 and contains five CpG sites, we performed BSP analysis of DNA isolated from two representative metastatic tissues and from SK-OV-3 cells. All five GABRP promoter CpGs were hypomethylated in the metastatic tissues compared with those in the SK-OV-3 cells, and the largest decrease in DNA methylation was observed at the −963 CpG site (Figure 3e).

GABRP expression was altered by DNA methylation-dependent epigenetic modifications

To determine whether GABRP is regulated by DNA methylation-dependent epigenetic modifications, we treated SK-OV-3 cells with the DNA methyltransferase inhibitor 5-aza-dc and then measured the levels of DNA methylation at the GABRP promoter CpGs by qMSP and of GABRP mRNA expression using RT-qPCR. Treatment with 5-aza-dc resulted in a decrease in DNA methylation at the −963, but not the −1142, CpG site (Figures 4a and b). Importantly, GABRP expression was increased ~1.8-fold, and in a dose-dependent manner, compared with that of untreated cells (Figure 4c). These data indicate that GABRP expression is regulated by a DNA methylation-mediated epigenetic mechanism, and the DNA methylation status of the −963 CpG site is critical for the transcriptional regulation of GABRP.
The GABRP promoter was hypomethylated in advanced stage, but not early-stage, primary ovarian tumors

To further validate the clinical significance of our findings, we next investigated the DNA methylation status at the −963 CpG site within the GABRP promoter, and GABRP expression in 15 paired clinical samples, which included primary ovarian carcinomas and adjacent normal tissues, from six patients diagnosed at early stage and nine patients at advanced stage. Although the DNA methylation at the −963 CpG site of the GABRP promoter was not significantly altered in early-stage primary tumor tissues, it was reduced in advanced-stage primary tumor tissues, compared with that of normal

Figure 2 GABRP promotes migration and invasion of SK-OV-3 cells through ERK pathway. SK-OV-3 cells were transfected with EGFP and GABRP expression constructs (a) or with siNC and siGABRP (b). After transfection, the effects on migration were determined using transwell assays, and the effects on invasion were determined using Matrigel-based invasion assays. Representative images of migrated or invaded cells are shown (×100 magnification). The amounts of migrated or invaded cells were quantified by measuring the absorbance of extracts from cell stains at 595 nm (a, b). After treatment with the MEK inhibitor U0129, the migration and invasion of SK-OV-3 cells were determined using transwell and Matrigel-based invasion assays, respectively (c). Immunoblot analysis of total ERK1/2 and phosphorylated ERK1/2 proteins was performed in cells treated with expression plasmid constructs, siRNAs or MEK inhibitor (d). A representative immunoblot of total ERK1/2 and phosphorylated ERK1/2 proteins (upper panel of d), and quantification of phosphorylated ERK1/2 relative to total ERK1/2 (lower panel of d), are shown. All data shown are means ± s.d.’s for triplicate measurements. Statistical analysis was performed with t-tests (*P<0.05, **P<0.01, ***P<0.001). siGABRP, GABRP siRNA; siNC, non-targeting control siRNA.

The GABRP promoter was hypomethylated in advanced stage, but not early-stage, primary ovarian tumors

To further validate the clinical significance of our findings, we next investigated the DNA methylation status at the −963 CpG site within the GABRP promoter, and GABRP expression in 15 paired clinical samples, which included primary ovarian carcinomas and adjacent normal tissues, from six patients diagnosed at early stage and nine patients at advanced stage. Although the DNA methylation at the −963 CpG site of the GABRP promoter was not significantly altered in early-stage primary tumor tissues, it was reduced in advanced-stage primary tumor tissues, compared with that of normal
tissues (Figures 5a and b). More specifically, 89% of the advanced-stage patients had DNA methylation levels at the −963 CpG site that were decreased by more than 20% in the tumor tissue relative to the paired normal tissue, whereas this decrease was only observed in 17% of the early-stage patients. Consistent with these data, the median value of relative GABRP expression was higher in advanced-stage tumor tissues than in early-stage tumor tissues, although, the data were not statistically significant (Figure 5c). To examine the association between the expression level of GABRP and aggressiveness of tumors in serous-type ovarian cancer, we performed microarray gene expression analysis using The Cancer Genome Atlas (TCGA) database, which include gene expression data from 138 patients with ovarian cancer and

Figure 3 DNA methylation at the GABRP promoter CpG sites was altered in metastatic tissues from mouse xenografts. The DNA methylation status of GABRP at the −1142 and −963 CpG sites was analyzed using the Illumina HumanMethylation 450 BeadChip (a, b) and qMSP (c, d). The error bars represent the means ± s.d.’s from three independent experiments. The DNA methylation status within the GABRP promoter region was analyzed by bisulfite sequencing (e). The GABRP promoter region is located between positions 170209520 and 170209971 of chromosome 5 in the human GRCh37/hg19 assembly and contains five CpG sites, which are located at −1178, −1142, −994, −963 and −924 from the transcription start site. Each circle represents a CpG dinucleotide. The methylation status of each CpG site is indicated with a black (methylated) or white (unmethylated) circle. The percentage of methylation at each site is indicated in a pie graph on the bottom line. The black segment of the pie graph is proportionally sized to represent the percentage of methylation. The triangles above the circles and bars indicate the CpG sites that were used for methylation array and qMSP analyses. The statistical analyses were performed using one-way ANOVA, followed by Bonferroni post tests (**P < 0.05). M, the percentage of methylated CpGs; U, the percentage of unmethylated CpGs.
lymphatic invasion and 80 patients with ovarian cancer but without lymphatic invasion. Regardless of lymphatic invasion, the average GABRP expression was higher in the advanced stage than in the early-stage tumor tissue, but the differences were not significant in either subgroup, with lymphatic invasion (P = 0.664) or without lymphatic invasion (P = 0.303) (Supplementary Table S2). These findings suggest that aberrant demethylation of specific CpG sites within the GABRP promoter is associated with the aggressiveness of ovarian carcinoma cells.

**DISCUSSION**

Although it is well known that GABA and GABA receptors function as predominant inhibitory neurotransmitters in the mature mammalian brain, their role in non-neuronal cells, including cancer cells, is largely unknown. More recently, it was reported that GABAergic signaling is involved in the development of many tissues, including the peripheral nervous system, lung, liver, pancreas and testis, and that it regulates proliferation, migration and invasion in different cell types. Several studies have also shown that GABA receptors are involved in tumorigenesis, migration, invasion and proliferation of cancerous cells, however, the functions of GABA receptors in these processes seem to be cancer-type dependent. For instance, Chen et al. reported that GABA-induced cell migration and invasion are suppressed in HCC human liver cancer cells through the GABAA receptor, whereas another study reported that the GABA A receptor alpha3 promotes breast cell migration by activating the AKT pathway.

The GABA receptor GABRP is overexpressed in pancreatic ductal adenocarcinoma cells and basal-like subtype of breast cancer, where it has been shown to promote proliferation, tumorigenesis and migration. Similarly, in this study, we showed that levels of GABRP were markedly upregulated in metastatic tissues from an ovarian cancer xenograft mouse model compared with those of SK-OV-3 cells. Furthermore, we performed GABRP gain- and loss-of-function studies in the SK-OV-3 ovarian carcinoma cell line to investigate the effect of GABRP on cell migration and invasion. The results showed that GABRP promoted cell migration and invasion of ovarian carcinoma cells, suggesting it has an essential role in the acquisition of an aggressive phenotype in ovarian cancer cells. Finally, treatment with the MEK inhibitor U0126 suppressed the migratory and invasive abilities of ovarian carcinoma cells,
indicating that these functions of GABRP are mediated via ERK pathway activation. Our newly identified function and mechanism of GABRP in ovarian cancer cells are consistent with those shown previously in pancreatic and breast cancer cells.

To address the mechanism of GABRP overexpression in metastatic ovarian cancer cells, we analyzed genome-wide DNA methylation profiling data and identified two GABRP promoter CpG sites at −1142 and −963 bp from the transcription start site that were hypomethylated in metastatic tissues from xenograft mice compared with those in the injected ovarian carcinoma cells. We further showed that treatment with a DNA methyltransferase inhibitor recovered the suppressed expression of GABRP, indicating that GABRP expression is regulated by a DNA methylation-dependent epigenetic mechanism, which had not been previously shown. Our results also showed that the DNA methylation status at the −963 CpG site was critical for the transcriptional regulation of GABRP.

To demonstrate the clinical relevance of our findings, we showed that the DNA methylation at the −963 CpG site of GABRP was decreased in primary ovarian tumors of advanced stage, but not of early stage, compared with that of their paired normal tissues, strongly suggesting that hypomethylation at the −963 CpG site is a marker for an aggressive form of ovarian cancer. Although not statistically significant, we also observed higher expression of GABRP in several primary tumors of advanced stage than in primary tumors of early stage. Hence, we speculate that ovarian carcinoma cells with upregulated GABRP expression, due to the hypomethylation of specific GABRP promoter CpG sites, acquire aggressive phenotypes, which subsequently leads to their metastasis.

CONFLICT OF INTEREST

The authors declare no conflict of interest.

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