IQGAP3 promotes cancer proliferation and metastasis in high-grade serous ovarian cancer

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Abstract. Ovarian cancer is a type of gynecological cancer with the highest mortality rate worldwide. Due to a lack of effective screening methods, most cases are diagnosed at later stages where the survival rates are poor. Thus, it is termed a ‘silent killer’ and is the most lethal of all the malignancies in women. IQ motif containing GTPase Activating Protein 3 (IQGAP3) is a member of the Rho family of GTPases, and plays a crucial role in the development and progression of several types of cancer. The aim of the present study was to investigate the oncogenic functions and mechanisms of IQGAP3 on the proliferation and metastasis of high-grade serous ovarian cancer (HGSOC). Therefore, the expression levels of IQGAP3 in HGSOC and normal tissue samples were compared, and IQGAP3 knockdown was performed to examine its functional role using various in vitro and in vivo experiments. It was demonstrated that the expression of IQGAP3 was upregulated in HGSOC tissues compared with the healthy tissues; this differential expression was also observed in the ovarian cancer cell lines. Functional experimental results suggested that IQGAP3 silencing significantly reduced proliferation, migration and invasion in ovarian cancer cell lines. Moreover, in vivo experimental findings validated the in vitro results, where the tumorigenic and metastatic capacities of IQGAP3-silenced cells were significantly lower in the nude mice compared with the mice implanted with the control cells. Furthermore, knockdown of IQGAP3 resulted in increased apoptosis, and the effects of IQGAP3 expression on various epithelial-mesenchymal transition markers were identified, suggesting a possible mechanism associated with the role of IQGAP3 in metastasis. The effect of IQGAP3 silencing on chemosensitivity towards olaparib was also assessed. Collectively, the present results indicated that IQGAP3 is a potential diagnostic and prognostic marker, and a putative therapeutic target of HGSOC.

Introduction

Ovarian cancer is one of the most malignant types of gynecological cancer, and is the 11th most common type of cancer among women, as well as the 5th leading cause of cancer-associated mortality in the USA (1). In addition, ovarian cancer is the leading cause of gynecological malignancy-associated mortality (1,2). The American Cancer Society estimated there were 22,530 new cases of ovarian cancer and 13,980 mortalities from ovarian cancer in the USA in 2019 (1). Furthermore, the incidence of new ovarian cancer cases has been decreasing on average by 2.5% each year in the past decade; however, the overall survival rate has not improved in recent years (2). The current 5-year survival rate for all the stages of ovarian cancer cases in the US is approximately 47% (1). However, approximately 60% of the new cases are diagnosed at advanced stages, and in those cases, the 5-year survival rate is only 29% (1). Moreover, there is a high rate of recurrence even after aggressive multimodal treatment, which further worsens the prognosis (3).

Ovarian cancer belongs to a group of heterogeneous tumors that arise spontaneously largely from the ovaries, but may evolve from various other potential sources (4-6). In addition, ovarian cancer can be morphologically classified into epithelial and non-epithelial types, of which 80-90% of all ovarian cancer cases are epithelial type (5). Based on their aggressiveness, epithelial ovarian cancers (EOCs) are further subdivided into high- and low-grade categories; or morphologically, they are subdivided into serous, endometrioid, mucinous and clear cell varieties (7,8). High-grade serous ovarian cancer (HGSOC) accounts for 50-60% of all ovarian neoplasms (8). Furthermore, advanced HGSOC accounts for approximately 50% of all EOCs (7-9), but the precise etiological factors underlying ovarian cancer have not been fully elucidated. However, hereditary susceptibility is considered an important risk factor, as approximately 35% of HGSOC cases harbor a germline mutation of the tumor suppressor genes Breast cancer type 1 susceptibility protein (BRCA1) or BRCA2 (10).
The current therapeutic measure used to treat ovarian cancer is a multimodal regimen, and a combination of platinum and paclitaxel is used as the primary chemotherapeutic regimen (11). However, the relapse rate remains high due to chemoresistance (12). Poly ADP-ribose polymerase inhibitor (PARPi) has been introduced as a promising therapeutic agent to improve the prognosis of HGSOC (13). Olaparib is the most commonly used PARPi, and exhibits favorable outcomes in lowering disease progression and mortality rates (14). Moreover, olaparib has been approved by the Food and Drug Administration (FDA) as the first monotherapy to combat advanced epithelial ovarian cancer cases harboring germline BRCA mutations (15).

IQ motif containing GTPase Activating Proteins (IQGAPs) are a family of GTPase activating proteins, which have been evolutionarily conserved from yeast to mammals (16,17). In a review by Hedman et al (18), the varied functions of IQGAPs, in addition to serving as scaffolding proteins are discussed. In total, three members of the IQGAP family have been described in humans (18). Furthermore, all three members are equipped with four IQ motifs and a Ras GTPase-activating protein (GAP)-related domain (18); the GAP-related domain of IQGAPs mediates its binding to the Rho family of GTPases (19). A member of the Rho family of GTPases, Cell Division Cycle 42 (CDC42) has been revealed to serve critical roles in cell proliferation, survival, adhesion and migration, and is correlated with a less favorable prognosis in various types of cancer (20-23). Of the three IQGAP family members, IQGAP1 has been reported to play a synergistic role in cancer progression and aid in cellular motility (24-26). However, IQGAP2 exhibits a tumor suppressive function (26). Moreover, IQGAP3 is hypothesized to be involved in the proliferation of epithelial cells (27), and is a novel member of the IQGAP family, which was discovered in 2007 (28). IQGAP3 is located on chromosome 1 at 1q21.3 loci and has been reported to act as an oncogene in several types of cancer (29-35). Furthermore, IQGAP3 is a transmembrane protein, and has been speculated to be a potential therapeutic target (35).

The present study aimed to analyze the differential expression of IQGAP3 in HGSOC and healthy tissues, and the effect of IQGAP3 knockdown on various functional processes, such as cell proliferation, migration, invasion and apoptosis, to determine whether IQGAP3 could serve as a potential oncogenic prognostic and therapeutic target for patients with HGSOC.

Materials and methods

**Tissue samples.** A total of 149 ovarian cancer tissue samples (patient age range, 34-79 years; median age, 56 years) and 64 healthy fallopian tube epithelial tissues (patient age range, 26-74 years; median age, 47 years) with detailed clinical information were collected from the Pathology Department at Qilu Hospital of Shandong University (Jinan, China) between January 2005 and January 2015. All the malignant samples were diagnosed in accordance with the International Federation of Gynecology and Obstetrics criteria (36). The healthy samples were collected from patients who underwent surgery for benign conditions. Signed consents were collected from all the patients and the study was approved by the Ethics Committee of Qilu Hospital of Shandong University.

Survival analysis was performed on datasets from the Gene Expression Omnibus (GEO) database, including 523 patients for overall survival analysis using datasets GSE18520 (37), GSE26193 (38), GSE30161 (39), GSE63885 (40) and GSE9891 (41), and 483 patients for the progression-free survival analysis using datasets GSE26193, GSE30161, GSE63885, GSE9891, GSE65986 (42) on Kaplan-Meier Plotter (43).

**Cell lines and cell culture.** Human ovarian cancer cells A2780 (cat. no. CL-0013; Procell Life Science & Technology Co., Ltd.) were cultured in RPMI-1640 medium supplemented with 10% FBS and penicillin (100 IU/ml) and streptomycin (100 µg/ml) (all Gibco; Thermo Fisher Scientific, Inc.). HEY cells (gifted from Dr Jianjun Wei; Laboratory at Northwestern University) were cultured in DMEM (Gibco; Thermo Fisher Scientific, Inc.) supplemented with 10% FBS. All the cells were maintained in a humidified incubator at 37°C with 5% CO₂.

**Immunohistochemistry (IHC).** IHC staining of the tissue microarray (TMA) was performed on 4-µm sections sliced from each TMA receiver block fixed with 4% paraformaldehyde at room temperature for 48 h and embedded in paraffin. Tissue slides were deparaffinized in xylene and rehydrated in a graded series of ethanol (10 min each in 100, 95, 80 and 70% ethanol). Antigen retrieval was performed using a heat-induced epitope retrieval method with 10 mmol/l EDTA buffer (pH 8.0) at 98°C for 15 min. Endogenous peroxidase activity was quenched with 3% hydrogen peroxide in methanol for 15 min at 37°C, and non-specific binding was blocked by incubation with donkey serum as part of the SP9000 IHC kit (OriGene Technologies, Inc.; cat. no. SP9000) for 30 min at 37°C. The slides were subsequently incubated overnight at 4°C in a humid chamber with anti-IQGAP3 (Abcam; cat. no. ab219354) antibody at a dilution of 3 µg/ml. Staining was visualized using I-View 3,3'-diaminobenzidine staining detection system (OriGene Technologies, Inc.; cat. no. ZLI-9018). The IHC score was determined using a semi-quantitative method based on the extent and intensity of positively stained cells. The percentage of positive cells within each sample was scored independently from 0 to 100% upon observation under a light microscope (magnification, x10). The intensity of immunostaining was graded as follows: 0, Negative; 1, weak; 2, moderate; and 3, strong. The final IHC score was generated by multiplying the percentage extent with the staining intensity score. Then, two gynecological pathologists independently reviewed the IHC staining. High IQGAP3 expression grade was defined as a final IHC score ≥100.

**Stable and transient transfection.** For stable transfection, lentiviral vector GV493 (hu6-MCS-CBh-gcGFP-IRES-puromycin) was packaged with IQGAP3 short hairpin (sh)RNA along with the respective negative control (NC), which were purchased from Shanghai GeneChem Co., Ltd. A total of 1x10⁵ cells were plated into 6-well plates 24 h prior to stable transfection. Multiplicity of infection (MOI) was determined and the lentivirus was added to the culture medium complemented with the transfection reagent HiTransGA (Shanghai...
GeneChem Co., Ltd.) with a MOI value of 20-50. After 24 h incubation, the medium was replaced with fresh culture medium containing 2 µg/ml puromycin (Sigma-Aldrich; Merck KGaA) for selection of the stably transfected colonies.

Transient transfection was performed using small interfering (si)RNAs purchased from Shanghai GenePharma Co., Ltd. at a concentration of 20 µM. RNAi-mediated knockdown was performed with the following siRNAs: si-IQGAP3-1, 5'-GGGAGGAGGAAGAAGGAU-3'; si-IQGAP3-2, 5'-GAGCAACCCAGGAGCUACUAU-3'; si-CDC42, 5'-GGAGCAGGAUUAUUCCCAACAGCGUGGACUGUTT-3'. Cells were transfected with Lipofectamine® 2000 (Invitrogen; Thermo Fisher Scientific, Inc.) according to the manufacturer's protocol. The subsequent experiments were performed 24-48 h after transfection.

**RNA extraction and reverse transcription-quantitative (RT-q) PCR.** Total RNA was extracted from tissue samples and cultured cells using TRIzol® reagent (Invitrogen; Thermo Fisher Scientific, Inc.) according to the manufacturer's protocol. mRNA was then reverse-transcribed into cDNA using PrimeScript cDNA Synthesis kit (Takara Bio, Inc.) at 37˚C for 1 h and then at 85˚C for 5 min according to the manufacturer's protocol. qPCR was performed using SYBR-Green Premix Ex Taq II (Takara Bio, Inc.) with a StepOne Plus RT PCR system (Applied Biosystems; Thermo Fisher Scientific, Inc.). The reaction conditions were as follows: Initial denaturation at 95˚C for 5 sec, followed by 40 cycles of annealing at 60˚C for 10 sec and an extension at 72˚C for 30 sec. β-actin was used as the endogenous control. The primers were designed based on the GeneBank sequences. The primer sequences used were: IQGAP3 forward, 5'-GGGAGGAGGAAGAAGGAU-3'; reverse, 5'-GGGAGGAGGAAGAAGGAU-3'; CDC42 forward, 5'-GGGAGGAGGAAGAAGGAU-3'; reverse, 5'-GGGAGGAGGAAGAAGGAU-3'; and β-actin forward, 5'-GGGAGGAGGAAGAAGGAU-3'; reverse, 5'-GGGAGGAGGAAGAAGGAU-3'. The relative gene expression level (44).

**Western blotting.** Cells were harvested and lysed in RIPA lysis buffer (Beyotime Institute of Biotechnology) with PMSF (1%) and NaF (1%). Protein samples were incubated for 30 min on ice and cell debris were removed by centrifugation at 12,000 x g at 4˚C for 15 min. The protein concentration was determined using a bicinchoninic acid (BCA) kit (Thermo Fisher Scientific, Inc.). Protein samples (30 µg) were separated by SDS-PAGE (5% stacking gel and 10% separating gel) and transferred to a PVDF membrane (EMD Millipore) using a Bio-Rad Trans-blot system (Bio-Rad Laboratories, Inc.). After blocking with 5% skimmed milk for 1 h at room temperature, the membrane was incubated overnight at 4˚C with the primary antibodies. The membranes were then rinsed with TBST (0.1% Tween-20) followed by incubation with a horseradish peroxidase-conjugated secondary antibody for 1 h at room temperature. Signals were detected using enhanced chemiluminescence (PerkinElmer, Inc.) with ImageQuant LAS 4000 (GE Healthcare Life Sciences). β-actin was used as the endogenous control. Densitometry analysis was performed using ImageJ version 1.52 g (National Institutes of Health).

The antibodies used were: Rabbit anti-human IQGAP3 (1:1,000; Abcam; cat. no. ab219354), rabbit anti-human CDC42 (1:1,000; Affinity Biosciences; cat. no. DF6322), rabbit anti-human Zinc Finger E-Box Binding Homeobox 1 (ZEB-1; 1:1,000; CST Biological Reagents Co., Ltd.; cat. no. 3396), rabbit anti-human N-cadherin (N-CAD; 1:1,000; CST Biological Reagents Co., Ltd.; cat. no. 13116), rabbit anti-human E-cadherin (E-CAD; 1:1,000; CST Biological Reagents Co., Ltd.; cat. no. 3195), rabbit anti-human Vimentin (1:1,000; CST Biological Reagents Co., Ltd.; cat. no. 5741), rabbit anti-human Snail (1:1,000; CST Biological Reagents Co., Ltd.; cat. no. 3879), rabbit anti phospho-(p-)AKT (pSer473) (1:1,000; Abcam; cat. no. ab66138), rabbit anti-human AKT (1:1,000; Abcam; cat. no. ab79463), rabbit anti-human PI3K (1:1,000; Abcam; ab182651), rabbit anti-human phosphorylated (p-)mTOR (1:1,000; CST Biological Reagents Co., Ltd.; cat. no. 2971), rabbit anti-human mTOR (1:1,000; CST Biological Reagents Co., Ltd.; cat. no. 2983), rabbit anti-human Bcl2 (1:1,000; CST Biological Reagents Co., Ltd.; cat. no. 2876), rabbit anti-human caspase3 (1:1,000; CST Biological Reagents Co., Ltd.; cat. no. 8G10), rabbit anti-human p-ATM Serine/Threonine Kinase (1:1,000; CST Biological Reagents Co., Ltd.; cat. no. 5883), rabbit anti-human AT1 (1:1,000; CST Biological Reagents Co., Ltd.; cat. no. 2873), rabbit anti-human Checkpoint Kinase 2 (CHK2; 1:1,000; CST Biological Reagents Co., Ltd.; cat. no. 2662), rabbit anti-human RAD51 (1:10,000; Abcam; cat. no. ab 33534) mouse anti-human Bax (1:1,000; CST Biological Reagents Co., Ltd.; cat. no. 2772), mouse anti-human caspase9 (1:1,000; CST Biological Reagents Co., Ltd.; cat. no. 9508), rabbit anti-human Pgp (1:1,000; Abcam; cat. no. ab 0347) and mouse anti-human β-actin (1:1,000; CST Biological Reagents Co., Ltd.; cat. no. 3700).

The secondary antibodies used were: Horseradish peroxidase-conjugated anti-rabbit (1:5,000; Sigma-Aldrich; Merck KGaA) or anti-mouse secondary antibody (1:5,000; Sigma-Aldrich; Merck KGaA; cat. no. A0545) or anti-mouse secondary antibody (1:5,000; Sigma-Aldrich; Merck KGaA; cat. no. A9044).

**Cell proliferation assay.** The proliferative ability of cells was measured using an MTT assay. Each cell line was seeded in quintuplicate into 96-well plates (0.8-1x10⁴ cells/well) for 0-4 days. At specified time points, 20 µl MTT reagent (Sigma-Aldrich; Merck KGaA) at 5 mg/ml concentration was added to each well, and the cells were incubated for an additional 3.5 h at 37˚C. Subsequently, the supernatants were discarded and 100 µl DMSO (Sigma-Aldrich; Merck KGaA) was added to each well. The absorbance at 490 nm was measured using a Varioskan Flash microplate reader (Thermo Fisher Scientific, Inc.).

**Cell migration and invasion assay.** Cell migration and invasion were analyzed using Boyden chamber-style cell culture inserts, with and without Matrigel (BD Bisciences), respectively. Matrigel was thawed at 4˚C and then coated onto the Transwell inserts, after which the gel was allowed to set at 37˚C for 1 h. Ovarian cancer cells (2x10⁵ cells) were seeded in the upper chamber of the Transwell inserts (24-well plate; 8-µm pore size; BD Bisciences) with 200 µl serum-free media. The lower chambers were filled with 700 µl culture media containing 10% FBS as the chemoattractant. After 6-48 h of incubation, the cells on the lower surface of the membrane were washed with PBS and fixed in 100% methanol for 15 min at room temperature. Then, cells were stained with...
0.1% crystal violet for 20 min at room temperature to quantify migration and invasion. Transwell inserts were observed under a light microscope (magnification, x10) and cells in 10 random fields were counted.

**Clonogenic assay.** For the colony formation assay, 500 cells were seeded into each well of a 6-well plate and maintained in media containing 10% FBS at optimum conditions of 37°C with 5% CO₂ for 10-14 days, until the colonies became visible to the naked eye. Colonies were then fixed with 100% methanol at room temperature for 15 min and stained with 0.1% crystal violet at room temperature. Colonies with >50 cells were counted manually under a light microscope (magnification, x10) for quantification.

**Apoptosis assay.** Apoptosis was detected using an Annexin V-FITC and propidium iodide (PI) kit (BD Biosciences), according to the manufacturer's protocol. A2780 and HEY cells were transfected with 20 µM si-IQGAP3 or si-NC, and were harvested 48 h after transfection with EDTA-free trypsin, centrifuged at 800 x g for 5 min at room temperature, washed twice with cold PBS, resuspended at a concentration of 1x10⁶ cells/ml and mixed with 100 µl 1X binding buffer. Subsequently, cells were stained with 5 µl Annexin V-FITC and 5 µl PI at room temperature for 15 min in the dark, after which 300 µl 1X binding buffer was added and the cells were analyzed by flow cytometry (FACSCalibur; BD Biosciences) within 1 h. The results were analyzed using FlowJo software version X.0.7 (FlowJo, LLC).

**Cell viability assay.** A total of 2x10⁵ cells/well were seeded in 96-well plates. The A2780 and HEY cells were exposed to olaparib (Selleck Chemicals; cat. no. AZD2281) at various final concentrations (0, 5, 10, 20, 40 and 80 µmol/ml) at 37°C for 36-72 h. Each concentration was repeated in quintuplicate wells. Subsequently, 20 µl 5 mg/ml MTT was added to each well. After incubation for 3.5 h, the medium was replaced with 100 µl DMSO, and cell viability was determined by analyzing the absorbance values at 490 nm on a Varioskan Flash microplate reader (Thermo Fisher Scientific, Inc.).

**Mouse xenograft models.** HEY cells that were stably transfected with IQGAP3-shRNA and the corresponding NC were used for the *in vivo* experiments. For *in vivo* experiments, eight female athymic BALB-c nude mice (age, 5 weeks; weight, 20-30 g) were purchased from Nanjing Biochemical Research Institute and housed in a standard pathogen-free condition in individually ventilated cages with HEPA filters at the ambient temperature of 30-31°C and humidity of 50-60% with 12 h light/dark cycle, and adequate access to food and water. For tumor formation assays, 1x10⁶ cells (knockdown or control), resuspended in 200 µl PBS were subcutaneously injected into either side of the axilla.

For metastasis assays, 1x10⁷ cells were intraperitoneally injected individually in the experimental and control groups. After 2-3 weeks, bioluminescence images were captured on an In-vivo imaging system (Kodak 2000 Imager). The mice were euthanized via intraperitoneal injection of 200 mg/kg sodium phenobarbital and the tumors were excised, fixed with 4% paraformaldehyde at room temperature for 48 h, paraffin-embedded and sectioned into 5-µm slices for hematoxylin and eosin staining. The tissue slides were stained with hematoxylin for 5 min and eosin for 10 min at room temperature and observed under a light microscope (magnification, x4).

**Statistical analysis.** GraphPad Prism version 7 (GraphPad Software, Inc.) was used to analyze data. A χ² test was used to analyze the differences in clinical characteristics. Survival analysis was performed using Kaplan-Meier analysis and a log-rank test. An unpaired Student's t-test and a one-way ANOVA were used to determine the statistically significant differences between different groups. Fisher's least significant difference was used for the post-hoc test following ANOVA. Data are presented as the mean ± standard deviation of ≥3 independent experiments. P<0.05 was considered to indicate a statistically significant difference.

**Results**

**IQGAP3 expression is upregulated in HGSOC.** The mRNA and protein expression levels of IQGAP3 in healthy fallopian tube and HGSOC tissues were determined using RT-qPCR and western blotting, respectively. The mRNA expression level of IQGAP3 was significantly higher in HGSOC tissues compared with the control samples (Fig. 1D). Furthermore, IQGAP3 protein expression was significantly upregulated in HGSOC tissues compared with the fallopian tubal samples (Fig. 1C).

**Upregulated expression of IQGAP3 is associated with a less favorable prognosis.** To examine whether upregulated expression of IQGAP3 was associated with clinical prognosis, IHC staining was performed on 149 HGSOC samples (Fig. 1E). Most positive staining was observed in the cytoplasm and at the cell membrane. Moreover, a high expression of IQGAP3 was observed in 53.02% (79/149) of tissues. Subsequently, the relationship between IQGAP3 and clinicopathological characteristics were assessed (Table I). The patients with a lower expression of IQGAP3 had longer survival times compared with those with higher IQGAP3 expression levels. A log-rank test demonstrated that the upregulated expression of IQGAP3 was significantly associated with overall survival (P=0.0149), as well as progression-free survival (P=0.0044; Fig. 1A).

Survival analysis performed on GEO cohorts using Kaplan-Meier Plotter, showed a significant association between IQGAP3 expression and both overall and progression-free survival (Fig. 1B). In addition, further analysis indicated that the expression of IQGAP3 was associated with several other clinicopathological parameters, including recurrence of the disease (P=0.0065), CA125 levels (P=0.0147) and peritoneal metastasis (P=0.0007; Table I).

**Downregulation of IQGAP3 reduces proliferation and colony formation of HGSOC ovarian cancer cells, and attenuates tumorigenicity in a xenograft model.** Downregulation of IQGAP3 resulted in the reduced proliferation of ovarian cancer cells *in vitro*. Moreover, two siRNAs, si-IQGAP3-1 and si-IQGAP3-2, were used to silence IQGAP3 in A2780 and HEY cells. MTT assays results identified a significant suppression of the proliferative capacity in the two cell lines.
following transfection with the siRNAs compared with the NCs (Fig. 2A).

These findings were further assessed in the in vivo experiments, where xenografts of BALB-c nude mice were established with injection of HEY cells stably transfected with sh-IQGAP3 or NC (Fig. 2C). After 3 weeks, the mice were euthanized, imaged on a bioluminescence imaging system, and the tumors were excised and weighed. It was found that...
IQGAP3 promotes proliferation and metastasis of ovarian cancer. To evaluate the role of IQGAP3 on metastasis of ovarian cancer in vivo, female nude mice were injected intraperitoneally with sh-IQGAP3-HEY cells or their corresponding NCs. Then, 3 weeks after injection, the mice were euthanized and the peritoneal cavities were examined for metastases. Consistent with the in vitro experimental results, mice injected with IQGAP3-silenced cells exhibited significantly lower numbers of metastatic nodules compared with the respective NC group (P<0.05; Fig. 3C). Bioluminescence imaging also identified larger metastatic foci in the control group compared with the knockdown group (Fig. 2E).

The excised metastatic nodules were fixed with formalin and paraffin embedded and 4-µm thick slices were sectioned. Subsequently, the slides were stained using hematoxylin and eosin staining (Fig. S1).

IQGAP3 knockdown promotes apoptosis in ovarian cancer cells. To assess the effects of IQGAP3 knockdown on apoptosis of ovarian cancer cells, Annexin V-FITC/PI dual staining was performed following transfection with si-IQGAP3 or NC. Both A2780 and HEY cells exhibited significantly increased apoptosis following knockdown of IQGAP3 compared with the respective NC group (Fig. 5A and B). These results were further validated by the increased expression of the pro-apoptotic proteins Bax, Caspase 3 and Caspase 9, and decreased expression of Bcl-2 following IQGAP3 knockdown (Fig. 4).

IQGAP3 knockdown increases sensitivity to chemotherapy with PARPi. The si-IQGAP3 transfected A2780 and HEY cells were exposed to various concentrations of olaparib (5, 10, 20, 40 or 80 µmol/ml) for 36-72 h, after which, the cell viability was assessed using an MTT assay. Cells transfected with si-IQGAP3 exhibited increased sensitivity to olaparib compared with the respective control group (Fig. 6A). Western blotting results identified the downregulation of the expression levels of Rad51, p-ATM (normalized to total ATM) and CHK2 when IQGAP3 expression was knocked down (Fig. 4). Thus, knockdown of IQGAP3 may have sensitized cells to olaparib by downregulating key factors involved in the DNA damage response.

Phosphoglycolate Phosphatase (Pgp) is a multidrug resistance protein that is localized in the cell membrane and is responsible for extruding several xenobiotics (including chemotherapeutic agents) outside the cells, rendering the cells chemoresistant (45). IQGAP3 knockdown reduced the expression of Pgp, which in turn attenuated chemoresistance (Fig. 4). Therefore, it was speculated that this effect may underlie the enhanced sensitivity of cells towards olaparib following IQGAP3 knockdown.

IQGAP3 exerts its function via the regulation of CDC42. It has been reported that IQGAP3 is an effector of CDC42 (28). Xu et al (35) also revealed that IQGAP3 may exert its oncogenic function in pancreatic cancer via the regulation of CDC42. To determine whether IQGAP3 was associated with CDC42 in ovarian cancer, the protein expression levels of CDC42 in ovarian cancer cells were assessed by knocking down IQGAP3 expression. It was identified that knockdown of IQGAP3 decreased the expression of CDC42 (Fig. 4).

Table I. Association between IQGAP3 expression and clinicopathologic characteristics.

<table>
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<th>Low expression</th>
<th>P-value</th>
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aP<0.05; bP<0.001. FIGO, International Federation of Gynecology and Obstetrics; CA125, cancer antigen 125.

There was a significant decrease in tumor size and tumor weight in the sh-IQGAP3 group compared with the NC group (Fig. 2D and E), supporting the in vitro results. Therefore, the results demonstrated the contribution of IQGAP3 to tumor proliferation.

Knockdown of IQGAP3 also significantly reduced colony formation in both A2780 and HEY cell lines (Fig. 2B).

IQGAP3 increases migration and invasion of ovarian cancer cells via epithelial-to-mesenchymal transition (EMT). Transwell assays were used to examine the role of IQGAP3 on migration and invasion in vitro. A2780 and HEY cells both had significantly decreased migratory and invasive capacities when IQGAP3 was knocked down compared with the respective NC (Fig. 3A and B).

Furthermore, the underlying mechanism contributing to this increase in tumorigenic features was determined by analyzing EMT-related factors. Knockdown of IQGAP3 had an effect on the expression of several EMT markers (Fig. 4). Silencing of IQGAP3 resulted in downregulation of mesenchymal markers, including ZEB-1, Vimentin, N-CAD and Snail, while the expression of the epithelial marker E-CAD was upregulated. Thus, it was suggested that IQGAP3 induced the migration and invasion of ovarian cancer cells via induction of EMT.
Figure 2. *IQGAP3* promotes the proliferation of HGSOC *in vitro* and *in vivo*. (A) MTT assay identified significantly reduced proliferation in A2780 and HEY cells following knockdown of *IQGAP3* using si-*IQGAP3*-1 and si-*IQGAP3*-2 compared with the corresponding control. (B) Silencing *IQGAP3* reduced colony formation in A2780 and HEY cells compared with the respective controls. (C) Knockdown of *IQGAP3* with sh-*IQGAP3* in HEY cell line. (D) Knockdown of *IQGAP3* resulted in reduced tumor forming capacity in nude xenograft mice compared with the corresponding controls. Control cells were injected in the right armpit and sh-*IQGAP3*-transfected cells were injected in the left armpit. The weight and the volume of the xenograft tumors were significantly different between the control group and the sh-*IQGAP3*-transfected group. (E) Bioluminescence imaging identified the effects of *IQGAP3* silencing on tumor formation, as well as metastasis *in vivo*. Control cells were injected in the left armpit and *IQGAP3* knockdown cells were injected in the right. Metastatic foci were more visible in the nude mice injected intraperitoneally with control (left) compared with sh-*IQGAP3*-transfected cells (right). *P<0.05; **P<0.01; ***P<0.001; ****P<0.0001 vs. NC.

*IQGAP3*, IQ motif containing GTPase Activating Protein 3; siRNA, small interfering; shRNA, short hairpin RNA; NC, negative control; shRNA, short hairpin RNA.
Therefore, the effects of CDC42 on the cancer cells were assessed. Knockdown of CDC42 expression (Fig. 6B) resulted in a significant decrease in the proliferative potential of HEY cells (Fig. 6C). Furthermore, migration and invasion were inhibited, while apoptosis was enhanced following CDC42 knockdown (Fig. 6D and E). Collectively, these results suggest that IQGAP3 may exert its effects via the regulation of CDC42.

Discussion

The principle dilemma when dealing with ovarian cancer is the rate of distant metastasis at the time of diagnosis and its resistance to chemotherapy, which frequently results in negative consequences (1,46-48). Thus, there is an unmet need for an improved understanding of the molecular mechanisms...
involved in the proliferation, metastasis and chemoresistance of ovarian cancer.

Out of the three primary members of the IQGAP family, IQGAP1 has been reported to be an oncogene, and IQGAP2 a tumor suppressor (24-26,49). Furthermore, IQGAP3 is a scaffolding protein, which interacts with various structural proteins that influence the cytoskeletal dynamics and intracellular signaling (28). IQGAP3 has also previously been implicated in the proliferation of epithelial cells (27). Moreover, previous studies have revealed the role of IQGAP3 in the proliferation and metastasis of lung, gastric, breast, pancreatic cancer, and colorectal cancer as well as hepatocellular carcinoma (29-35). Therefore, the role of IQGAP3 is crucial in the malignant transformation of several types of cancer. Yang et al (29) also reported that IQGAP3 promotes the metastasis of lung cancer cells by activating the epidermal growth factor receptor/ERK signaling pathway. In addition, Yang et al (29) used bioinformatics analysis to show that IQGAP3 is upregulated in several malignancies, including ovarian cancer. Furthermore, Wu et al (50) reported there were alterations in the genes regulating cytoskeleton remodeling in metastatic lung adenocarcinoma and that IQGAP3 was a marker of a less favorable prognosis.

The present results indicated that IQGAP3 was upregulated in ovarian cancer, and this enhanced expression resulted in increased proliferative and metastatic capacities in vitro and in vivo. Upon silencing IQGAP3, the aggressive nature of ovarian cancer cells was significantly abrogated. Thus, IQGAP3 may be a putative oncogene in HGSOC. Moreover, the upregulated expression of IQGAP3 was associated with a shorter overall and progression-free survival, cancer recurrence and CA125 expression. Kaplan-Meier survival analysis on data obtained from the online GEO database also demonstrated that patients with an upregulated expression of IQGAP3 exhibited reduced survival rates, further validating the in vitro and in vivo results. However, whether IQGAP3 is an independent poor prognostic factor of HGSOC is yet to be determined.

A number of in vitro and in vivo experiments were designed to establish the oncogenic potential of IQGAP3 in HGSOC. IQGAP3 expression was significantly upregulated in HGSOC compared with the healthy control. Cell proliferation
Figure 5. IQGAP3 increases apoptotic potential in cancer cell lines. Flow cytometry analysis indicated increased early, as well as late apoptosis in (A) A2780 and (B) HEY cells following knockdown of IQGAP3. ***p<0.001 vs. NC. IQGAP3, IQ motif containing GTPase Activating Protein 3; NC, negative control; si, small interfering RNA; PI, propidium iodide.
and tumorigenesis assays in nude mice demonstrated the decreased proliferative capacity of ovarian cancer cells when \textit{IQGAP3} was knocked down \textit{in vitro} and \textit{in vivo}.

Metastasis is a culmination of cancer cells gaining migratory and invasive abilities (51). Furthermore, distant metastasis at the time of diagnosis is one of the major obstacles negatively impacting the prognosis of ovarian cancer (52). The molecular mechanisms of metastasis in ovarian cancer are yet to be fully elucidated; however, EMT has been considered to be a potential contributing factor (52-54).

The present results suggested that \textit{IQGAP3} serves a substantial role in migration and invasion of ovarian cancer, and knocking down \textit{IQGAP3} reduced the metastatic potential. These findings were also observed \textit{in vivo}, where fewer metastatic foci formed in the mice injected with \textit{IQGAP3}-silenced cells compared with the control group.
EMT is initiated by several EMT-inducing transcription factors (53-58). In the present study, it was found that several of the EMT-inducing factors were affected by alterations in the expression of IQGAP3, which suggests a pivotal role of IQGAP3 in the induction of EMT in ovarian cancer.

Several studies have reported that the PI3K/AKT/mTOR signaling pathway is a crucial pathway by which cancer cells exhibit increased proliferative and metastatic potential (59-64). This signaling pathway is involved in several fundamental processes in ovarian cancer, such as cell proliferation, survival, autophagy, transcription regulation and angiogenesis (63,64). Therefore, to determine the mechanism underlying the effects of IQGAP3 in ovarian cancer, the effects of altering IQGAP3 gene expression of the PI3K/AKT/mTOR pathway were determined. Western blot analysis revealed a significant downregulation in the expression levels of PI3K, p-AKT and p-mTOR when IQGAP3 expression was knocked down. Thus, these results suggested that IQGAP3 may promote tumor progression and metastasis via the PI3K/AKT/mTOR signaling pathway.

Previous studies have shown that increased apoptosis may underlie decreased tumor growth, chemo-resistance and metastasis in several types of cancer (65,66). It has also been reported that IQGAP3 in certain types of cancer is closely associated with apoptosis (35). In the present study, it was demonstrated that the apoptotic potential of cells was increased when IQGAP3 expression was knocked down, and this may underlie the effects of IQGAP3 on tumor growth.

CDC42 is a member of the Rho family of GTPases, and is ubiquitously expressed (23). Moreover, CDC42 participates in the regulation of cytoskeletal dynamics, cellular proliferation, motility, polarity and cytokinesis (67). Wang et al (28) also identified a direct interaction between IQGAP3 with CDC42, and revealed IQGAP3 is an indispensable effector of CDC42-mediated cell proliferation. Furthermore, Xu et al (35) hypothesized that IQGAP3 may serve as an oncogene in pancreatic cancer by regulating the CDC42 signaling pathway. Morgan et al (68) reported there was an interaction between IQGAP3 and CDC42 using immunoprecipitation assays. The results of the present study also demonstrated that knocking down IQGAP3 expression resulted in the downregulation of CDC42 expression. Therefore, the role of CDC42 in ovarian cancer cells was further investigated. It was found that knockdown of CDC42 resulted in a significant decrease in proliferation, migration and invasion, and increased apoptosis in ovarian cancer cells. Thus, it was speculated that IQGAP3 may exert its function via modulation of CDC42, but further studies are required to verify this hypothesis.

PARP serves a key role in the DNA damage response of the cell (10,11,13). The PARP inhibitor olaparib has been recently approved by the FDA for the treatment of patients with ovarian cancer who harbor BRCA1/2 mutations (15). Moreover, BRCA1/2 mutations are responsible for 18-40% of lifetime risk ovarian cancer cases in women, and 5-15% of all diagnosed cases harbor one of these mutations (69). Thus, the introduction of PARPi, such as olaparib, may improve the prognostic prospects of patients. However, this drug is not effective for all HGSOc cases (70). IQGAP3 is associated with olaparib drug sensitivity, and knockdown of IQGAP3 in the present study resulted in increased efficacy of olaparib, suggesting that the effectiveness of the treatment may be dependent upon specific clinical aspects. Additionally, the expression profiles of proteins involved in DNA damage response of cell, including ATM and CHK, were assessed. In the present study, there was a significant decrease in the expression levels of these proteins following IQGAP3 knockdown. Furthermore, similar effects were observed in Rad51, which possesses a crucial role in the homologous recombination repair of DNA (10). Therefore, it was hypothesized that downregulation of DNA repair factors may result in defective DNA repair in cells, thus increasing the sensitivity to PARPi.

However, further investigations focusing on the mechanistic role of IQGAP3 in proliferation and metastasis of ovarian cancer are required before IQGAP3 may be considered a diagnostic and prognostic marker, and as a potential therapeutic target for ovarian cancer. To the best of our knowledge, the present study was the first to report the role of IQGAP3 in the progression of HGSOc.

In conclusion, IQGAP3 exhibited oncogenic features in HGSOc. In addition, the expression of IQGAP3 was upregulated in HGSOc, and its expression was associated with a poor outcome in patients. However, more studies are required to further validate IQGAP3 as a prognostic marker and a therapeutic target for ovarian cancer.

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Availability of data and materials

All data generated or analyzed during this study are included in this published article.

Authors’ contributions

SD, QZ and BK contributed to the conceptualization of the study. SD drafted the manuscript. SD, CQ, CS, ZZ and HW contributed to data acquisition, carried out the data analysis and revised the manuscript. QZ and BK were involved in analyzing the critical intellectual content and gave the final approval of the version to be published. All authors read and approved the final manuscript.

Ethics approval and consent to participate

Signed consents were collected from all the patients and the study was approved by Ethics Committee of Qilu hospital of Shandong University. Approval of Shandong University...
Animal Care and Use Committee was acquired for all the animal experiments.

Not applicable.

The authors declare that they have no competing interests.

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