Mechanism of resistin-induced enhancement of proliferation and migration of ovarian cancer cells

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Abstract

Introduction: Resistin, a novel hormone secreted by human adipocytes and mononuclear cells, is associated with obesity, insulin resistance, and inflammation. Recent studies showed that resistin plays a key role in ovarian cancer cells. In this study, we investigated the potential of resistin to regulate the proliferation and migration of ovarian cancer cells.

Material and methods: A series of in vitro functional experiments were carried out to elucidate the role of resistin in ovarian cancer progression and the molecular mechanisms underlying its role.

Results: Resistin enhanced the proliferation of human ovarian epithelial carcinoma cells (HO-8910) in a time- and dose-dependent manner (30–100 ng/ml). Furthermore, HO-8910 cells cultured in adipocyte-conditioned medium showed dramatically increased rates of proliferation. Resistin knockout during adipocyte culture attenuated the proliferation of HO-8910 cells treated with adipocyte-conditioned media, indicating that resistin may promote HO-8910 cell proliferation via the mechanistic target of a rapamycin (mTOR)-mediated signaling pathway. Resistin (30–100 ng/ml) also enhanced wound-healing rates in a time- and concentration-dependent manner. Co-culturing HO-8910 cells with adipocytes also increased the wound-healing rates. Resistin expression was inhibited by miR-124-1 transcriptional activity, and resistin-mediated HO-8910 cell migration was also regulated by miR-124-1. Furthermore, we also confirmed the role of resistin in promoting tumor growth in vivo.

Conclusions: These findings suggest that resistin may serve as an effective therapeutic target for ovarian epithelial carcinoma, especially in patients who are obese.

Key words: obesity, adipocyte, mTOR, miR-124.

Introduction

Ovarian cancer is the second most common gynecologic cancer worldwide. As the ovaries are located in the pelvic cavity and the peritoneum, early-stage ovarian cancer patients are generally asymptomatic. In approximately 70% of patients, ovarian cancer causes pathological changes beyond the ovaries. The autocrine secretion/self-secretion hypothesis proposes that the activation of oncogenes, such as that in tumor cells, can enable the production of certain growth factors within these tumor cells; this, in turn, promotes tumor proliferation and migration [1]. To develop therapeutic targets, we investigated the mecha-
nisms involved in the proliferation and metastasis of ovarian cancer. Resistin is a relatively new cytokine that was discovered in 2001 and plays a role in insulin resistance [2]. High levels of resistin are associated with the development of breast, gastric, colorectal, and esophageal cancers [3–6]. Increased resistin levels are associated with endometrial carcinoma, polycystic ovary syndrome, and uterine cancer [7]. Our preliminary studies indicated that increased resistin expression contributes to the proliferation and migration of ovarian cancer cells; however, this has not been confirmed in any other reported literature in China or abroad. In our present study, we investigated the role of resistin in the proliferation and migration of ovarian cancer cells. Determining the mechanisms underlying resistin actions will promote the development of targeted therapeutics for patients with ovarian cancer.

Material and methods

Materials

The HO-8910 human ovarian epithelial carcinoma cell line was purchased from the American Type Culture Collection (ATCC, Manassas, VA, USA). Roswell Park Memorial Institute (RPMI) 1640 medium was purchased from HyClone Co. (Logan, UT, USA). Recombinant resistin was purchased from Phoenix Pharmaceuticals (Burlingame, CA, USA). siRNA (small interfering RNA) was synthesized by Invitrogen using a block-it RNAi design. The JetPEI reagent was purchased from Polyplus-transfection SA (Illkirch, France). All other chemicals and drugs were purchased from Sigma Chemical (St. Louis, MO, USA). Female BALB/c nude mice (4 weeks of age) were obtained from Beijing Huafukang Biotechnology Co. Inc (Beijing, China).

Cell culture

HO-8910 cells were cultured in RPMI 1640 containing 10% fetal bovine serum (FBS) and penicillin/streptomycin (100 U/ml) in a humidified 37°C incubator. When confluent, cells were treated with resistin (30–100 ng/ml) for 6 to 48 h. For co-cultured experiments, HO-8910 cells were co-cultured with adipocytes and then treated with resistin at 100 ng/ml for 12 h.

MTT assay

Briefly, HO-8910 cells were seeded at a density of $1 \times 10^4$ cells/μl in a 96-well plate containing DMEM (Dulbecco’s modified Eagle medium) and incubated with varying concentrations (0, 10, 30, 100 ng/ml) of resistin for 12, 24, or 48 h at 37°C. Then, 100 μl aliquots containing MTT (3-(4,5-di-methylthiazol-2-yl)-2,5-diphenyl tetrazolium bromide; 5 mg/ml in complete DMEM) were added, and cells were incubated for an additional 5 h. MTT-containing medium was removed and 150 μl of dimethyl sulfoxide (DMSO) was added to dissolve the formazan crystals formed by live cells. Absorbance was measured at 570 nm, and cell viability (%) was calculated according to the following equation.

Isolation and primary culture of rat adipocytes

The experimental setup and all animal handling procedures were conducted in accordance with the Animal Handling Guidelines of Shengjing Hospital of China Medical University. Thirty male Wistar rats (Shengjing Hospital Laboratories, Shenyang, China), 6 weeks old, weighing 160–180 g, were housed in two-rat cages with standard bedding material at 21–22°C and 50–60% relative humidity with a 12-h light/dark cycle. Standard rat chow and water were provided ad libitum. After 2 weeks of acclimatization, the rats were sacrificed under isoflurane anesthesia via exsanguination from the exposed aorta using dry heparinized syringes and then rapidly decapitated. Following this, the fat pads were minced into 1 mm³ fragments and transferred to 15 ml centrifuge tubes containing 100 μl of collagenase I working solution. The tissues were then incubated in a 37°C water bath and agitated at 120 rpm for 50 min. After digestion, the suspensions were filtered through a 250 μm mesh filter into tubes containing 5 ml of preheated DMEM per tube. The filtered crude suspension of isolated cells was allowed to stand for 5 min in stopped polypropylene syringes at 22–24°C. The adipocytes floated to form a defined upper layer. Next, the lower aqueous fraction was slowly drained off, and syringes were recapped to retain the adipocytes. The adipocytes were washed three times using this technique. The final supernatant fraction contained intact adipocytes and a free fat layer resulting from broken adipocytes. Working adipocyte samples were acquired from the central layer of adipocytes, ensuring that the free fat layer was excluded.

Wound-healing assay

HO-8910 cells were seeded onto 24-well plates (at a density of $5 \times 10^4$ cells/well) containing RPMI 1640 medium supplemented with 10% FBS. The cells were incubated for 24 h until they reached 100% confluence or had overgrown. A sterile 10 μl pipette tip was used to make a scratch (wound) in the center of the wells, after which the wells were washed with phosphate-buffered saline (PBS). Following this, media containing different concentrations of resistin (0, 10, 30, and 100 ng/ml)
were added to the wells. The culture media were aspirated at 2, 4, 6, and 12 h, washed three times with PBS, and the healing of the wounded area was observed under a microscope. The healed area was evaluated using ImageJ software, and the rate of healing was calculated using the formula: \( \text{rate of healing} = \frac{D_2 - D_1 \times 100}{D_1} \), where \( D_1 \) represents the wounded area at 0 h after the scratch, and \( D_2 \) represents the wounded area after 12 h of healing. The experiment was repeated three times.

**Synthesis and transfection of resistin siRNA**

Resistin siRNA (Sense: 5’-GGACGUCCGUGAGGUCCUCACUGU-3’; Antisense: 5’-ACACAUGUUAUCUCAGGACGUCC-3’; each strand with a 3’ dTdT overhang) and negative control siRNA were purchased from Invitrogen Life Technologies. siRNA transfection was performed using Lipofectamine RNAiMAX and Opti-MEM (both from Invitrogen Life Technologies) according to the manufacturer’s instructions. On day 2, HO-8910 cells were incubated with 50 nM of each siRNA in a maturation medium and cultured until day 5.

**Plasmid construction and transient transfection assays**

Next, we constructed the fusion gene consisting of human miR-124 promoter and luciferase reporter. First, we purified a series of fragments (positions -1754 to +28) of human genomic DNA from peripheral blood, inserted them into pGL3 Basic Luciferase Reporter Vector (Promega, Madison, WI, USA), and designated them as p-miR124. HO-8910 cells were plated in 24-well plates, transfected with various plasmids via a jetPEI reagent, and incubated for 24 h. Thereafter, the cells were harvested and homogenized. DNA transfection was performed with 0.5 μg of fusion reporter genes and 10 ng of the internal reference gene pRL-TK (thymidine kinase promoter – Renilla luciferase reporter; Promega). Luciferase activity was demonstrated in rat adipocytes. Real-time quantitative PCR and western blotting analyses showed that resistin levels in adipocytes’ mRNA were reduced to 38% (Figure 1 B) and the protein levels were reduced to 32% (Figure 1 C).

**Effects of differently conditioned media on HO-8910 cell proliferation**

HO-8910 cells were incubated with cultured primary adipose cells, cos-7 cells (used as negative controls), or with harvested adipocyte-conditioned media. The MTT assay results showed that differently conditioned media can enhance the proliferation rate of HO-8910 cells at a given dose within 24 h, compared to cos-7 cell-conditioned medium, which can augment proliferation rates by 48 h (Figure 1 D; \( *p < 0.05, **p < 0.01 \)). Specific knockdown of resistin during adipocyte culture significantly reduced the proliferation-inducing ability of adipocyte-conditioned media. Our results show that adipocyte-conditioned media could significantly promote the proliferation rate of HO-8910 cells, indicating that adipokines secreted by adipose tissue may promote the proliferation of HO-8910 cells, particularly in obese patients.
Although the receptor of resistin has not been identified, mTOR (mechanistic target of rapamycin) is associated with multiple cell-surface receptor types and is implicated in cellular proliferation. To investigate whether mTOR activation was involved in the resistin-mediated response, we used western blotting to detect the phosphorylation of mTOR and its downstream target protein S6 in HO-8910 cells. Stimulation with resistin significantly elevated the phosphorylation levels of mTOR and S6 (Figure 2 A). Rapamycin is a macrolide immunosuppressant that inhibits mTOR protein kinase. HO-8910 cells were treated with rapamycin (10, 20, and 30 nM) and incubated for 24 h. The MTT assay revealed that rapamycin can significantly inhibit HO-8910 cell proliferation rates (control group: *p < 0.05; resistin-treated cells group: *p < 0.01) (Figure 2 B). Thus, resistin may promote HO-8910 cell proliferation via the mTOR signaling pathway.

Effect of resistin on HO-8910 cell migration

The wound-healing assay results showed that resistin promoted HO-8910 cell migration. Resistin (100 ng/ml) induced cell healing in a time- and dose-dependent manner. As early as 6 h after wound induction, the healing rate increased to 42.59 ±3.94% and remained elevated for up to 12 h (66.61 ±4.69%) (Figures 3 A, B). Treatment with resistin (30–100 ng/ml) for 12 h also caused a concentration-dependent increase in the wound-healing rate (43.31 ±1.28%, 53.65 ±6.01%, 66.61 ±4.69%) (Figures 3 C, D) (*p < 0.05, **p < 0.05).

The wound-healing assay suggested that the migration of HO-8910 cells co-cultured with resistin...
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resistin-pretreated adipocytes was enhanced, exhibiting a wound-healing rate of 44.88 ±3.83%. Resistin knockdown in the co-culture system significantly reduced the wound-healing rate (30.91 ±3.21%; *p < 0.05) (Figures 3 E, F). HO-8910 cells were then incubated with cultured primary adipocytes, cos-7 cells (used as negative controls), or harvested adipocyte-conditioned media. The adipocyte-conditioned media significantly enhanced the healing rate of HO-8910 cells compared to that induced by cos-7 cell-conditioned media (48.39 ±2.88%; *p < 0.05). Similarly, resistin knockdown in the co-culture system significantly reduced the wound-healing rate induced by adipocyte-conditioned media (36.33 ±2.01%; *p < 0.05) (Figures 3 G, H).

**Effects of miR-124 on resistin-treated HO-8910 cell migration**

We examined the effect of resistin on miR-124 expression and found that resistin significantly inhibited the expression of miR-124-1, but not that of miR-124-2 and miR-124-3 (Figure 4 A). Next, we cloned the gene sequence of miR-124-1 and subcloned it into the pGL3 Basic Luciferase Reporter Vector. We then measured the relative luciferase activity in the transiently transfected HO-8910 cells. Our results indicated that resistin treatment significantly inhibited the activity of miR-124-1 in HO-8910 cells (Figure 4 B). The exact mechanisms of the effects of miR-124 on migration must be further elucidated and will be addressed in our future studies.

**Resistin promotes tumor growth in vivo**

HO-8910 cells with sh-driven resistin knockdown were used to model the in vivo effects of resistin on the growth of tumors in nude mice. The results showed that tumors in the mice from the sh-resistin group, which were subcutaneously implanted with resistin-deficient HO-8910 cells, formed later, grew slower, and had significantly smaller volumes and weights compared with those in mice from the shRNA-NC and the control groups (*p < 0.05; Figure 5).

**Discussion**

This study’s results demonstrated that treatment with resistin promoted HO-8910 cell proliferation via the mTOR pathway and migration via the miR-124-1 pathway. Resistin upregulated the proliferation of HO-8910 cells in a time- and concentration-dependent manner. Knockdown of resistin expression via siRNA treatment reduces the proliferation rate of HO-8910 cells. This resistin-mediated increase in HO-8910 cell proliferation occurred via the mTOR signaling pathway, possibly through a paracrine/self-secretion effect. Resistin also upregulated the wound-healing rate in HO-8910 cells in a time- and concentration-dependent manner and reduced the activity of the miR-124-1 promoter. To the best of our knowledge, this is the first report demonstrating resistin-mediated signaling pathways involved in the promotion of proliferation and migration of ovarian cancer cells.

Ovarian cancer, one of the five types of malignancy observed in the female reproductive system, is often diagnosed too late for effective treatment. Although the diagnostic rate of ovarian cancer has been improving with advances in surgical techniques, chemotherapy, and targeted therapy, the spread of ovarian cancer to other organs is a key problem that remains to be solved. A study conducted in the United States showed that women with a normal weight experience lower mortality rates than those with obesity [8]. Central or ab-

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**Figure 2.** Involvement of mTOR (mechanistic target of rapamycin) signaling pathway in resistin-mediated induction of HO-8910 proliferation. A – Stimulation with resistin significantly elevated the phosphorylation of mTOR and S6 in HO-8910 cells. B – The mTOR inhibitor rapamycin significantly inhibited proliferation of HO-8910 cells that were pretreated with resistin.
Figure 3. Effect of resistin on HO-8910 cell migration and interference of siRNA in the resistin-mediated effect on co-culture migration. A, B – Time course of the effects of treatment with 100 ng/ml resistin on the wound-healing rates of HO-8910 cells. C, D – Dose-response effect of 12 h treatment with resistin on the wound-healing rate of HO-8910 cells. E, F – The wound-healing rate with resistin compared to that with the knockout of resistin in the co-culture system. Co-cultured cells were incubated with 100 ng/ml resistin for 12 h. G, H – Cos-7 cells were used as negative controls; adipocyte-conditioned media significantly promoted the wound-healing rate of HO-8910 cells. The wound-healing rate was attenuated by the knockout of resistin in a co-culture system.
Abdominal obesity is gaining popularity as a key risk factor for cardiovascular disease (CVD) and higher mortality rates [9, 10]. Obese patients show a low survival rate and a high risk of cancer recurrence, regardless of the administration of chemotherapy or other auxiliary therapies. However, the role of obesity in tumor development is unclear. Obesity causes a chronic, low-grade inflammatory state in which high levels of inflammatory adipokines are secreted by adipocytes. Adipocyte factors such as leptin, tumor necrosis factor α, interleukin-6, and adiponectin are closely associated with tumor development, especially in women with endometrial and breast cancer [11, 12]. Adipocytes can also
Furthermore, adipose tissue can secrete hormones or other types of proteins that promote the excessive growth of ovarian cancer cells. Previous studies have shown that obesity impacts ovarian cancer-induced mortality by influencing tumor biology; however, the relationship between ovarian cancer-induced mortality and obesity needs further research.

Resistin is an adipocytokine discovered in 2001 and named for its role in insulin resistance [2]. Current studies on resistin have mainly examined its roles in inflammation and insulin resistance. When insulin levels are constant, resistin can stimulate glucose output. Mice with hereditary and diet-induced obesity demonstrate significantly increased plasma resistin levels. In humans, resistin is not only secreted by adipocytes, but also expressed by peripheral blood monocytes [14]. Epidemiological studies show that plasma resistin levels are positively correlated with BMI. High levels of resistin are associated with the development of breast, gastric, colorectal, and esophageal cancer. High resistin expression is also associated with more malignant clinicopathological statuses and adverse prognoses [15]. Regarding gynecological diseases, studies have focused on the role of resistin in polycystic ovary syndrome. Increased resistin expression is also closely related to the development of choriocarcinoma and endometrial carcinoma. However, few studies have examined the role of resistin in ovarian cancer. Recently, Qiu et al. reported that increased resistin expression increases growth, invasiveness, and cisplatin resistance in ovarian cancer cells. The mechanisms driving this activity include the induction of epithelial-mesenchymal transition (EMT) and stemness, as evidenced by the downregulated expression of the epithelial marker e-cadherin and the upregulated expression of mesenchymal markers vimentin/ZEB1 and stemness markers sox2, oct4, and nanog [16]. Previously, we demonstrated that resistin promotes the angiogenesis of ovarian cancer cells [17]. This study showed that resistin promoted the proliferation of epithelial ovarian HO-8910 cancer cells. Knockdown of resistin in adipocyte cultures significantly decreased the proliferation of HO-8910 cells that were treated with the media used in th

mTOR is a key cellular molecule that determines the energy levels inside and outside the cell, regulates transcription and translation, and mediates cell and tissue responses during survival, growth, proliferation, and differentiation. Whereas mTOR inhibits the activity of its downstream target protein S6, rapamycin inhibits the activity of anabolic S6 ribosomal proteins and increases the transcription of proliferating cell nuclear antigen (PCNA), thereby regulating the cell cycle and inhibiting the proliferation of smooth muscle and endothelial cells and fibroblasts. Rapamycin also inhibits the transcription of proto-oncogene Bcl-2, which plays a key role in the cell cycle. Inhibition of the PI3K/AKT/mTOR signaling pathway is deemed a striking opportunity for the development of novel anticancer agents [18].

Our current study shows that resistin promoted the migration of epithelial ovarian cancer cells (HO-8910) as observed in the wound assay. Cell migration was attenuated by knocking down resistin expression in HO-8910 cells. Conversely, co-culturing ovarian cancer cells and adipocytes significantly promoted the migration of HO-8910 cells. Cell migration was attenuated by knocking down resistin expression in co-cultured cells. These results further showed that resistin plays an important role in the development and progression of ovarian cancer.

Similar to protein-encoding genes, miRNA is involved in important biological functions, and numerous miRNAs play vital roles in tumorigenesis and tumor inhibition. Tumorigenesis-associated miRNAs tend to promote cell proliferation, inhibit apoptosis and activity of immune cells, and control cell cycles. Suppressor miRNAs tend to inhibit cell growth and promote apoptosis. Multiple factors are involved in the migration and invasion of tumor cells. Recent studies have shown that endogenous non-coding small-molecule RNAs (microRNAs) play important roles in regulating gene transcription and protein translation. Among these miRNAs, miR-124 helps inhibit tumor invasiveness [19]. Studies show that miR-124 expression is downregulated in liver and cervical cancer at the cellular and tissue level, which inhibits the growth and infiltration of myoblastoma and oral
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Squamous cell carcinoma. miR-124 may act as a potential tumor suppressor and is known to exert important regulatory effects on cell migration during embryonic stem cell differentiation. In our present study, resistin significantly inhibited the expression of miR-124. Furthermore, observations from in vivo tumorigenesis experiments in nude mice also verified that the expression of resistin was positively correlated with the tumorigenic capacity of the cells. Our results suggest that resistin may reverse the inhibitory effects of miR-124, thereby promoting the migration ability of ovarian cancer cells. The validation of this notion, however, requires further studies.

In conclusion, the results of our present study suggest that targeting resistin, mTOR, and miR-124 expression may be a beneficial strategy for the treatment of patients with ovarian epithelial carcinoma, particularly in those with concurrent obesity. We found that resistin is involved in the progression of ovarian cancer and in proliferation and migration of ovarian cancer cells. Therefore, the mechanisms involved in resistin-mediated signaling may present new therapeutic targets in halting the development and migration of ovarian cancer. Our study also illustrates a new mechanism that promotes the migratory ability of ovarian cancer cells. Resistin promotes the proliferation of these cells via the mTOR pathway and their migration via regulating miR-124 activity. These results show that mTOR and miR-124 can be used as therapeutic targets in the treatment and prevention of metastasis in patients with ovarian cancer. We focused on the ability of resistin to promote the proliferation and invasion of ovarian cancer, and have only elucidated one of the potential pathways. Further research is required to uncover the other pathways that govern resistin-mediated effects in ovarian cancer, so that therapeutic targeting of resistin may be achieved in obese patients with ovarian cancer.

Conflict of interest

The authors declare no conflict of interest.

References