

AGR2, a mucinous ovarian cancer marker, promotes cell proliferation and migration

Kyoungsook Park¹, Yong Jin Chung¹,
Hyekyung So¹, Kwangsoo Kim¹, Junsoo Park⁴,
Mijoung Oh¹, Minwha Jo¹, Kyusam Choi¹,
Eun-Ju Lee⁵, Yoon-La Choi², Sang Yong Song²,
Duk-Soo Bae³, Byoung-Gie Kim³ and Je-Ho Lee^{2,6}

¹Samsung Biomedical Research Institute

²Department of Pathology

³Department of Obstetrics and Gynecology
Samsung Medical Center

Sungkyunkwan University School of Medicine
Seoul 135-710, Korea

⁴Division of Biological Science and Technology
Yonsei University

Wonju 220-710, Korea

⁵Department of Obstetrics and Gynecology
Chung-Ang University School of Medicine

Seoul 156-755, Korea

⁶Corresponding author: Tel, 82-2-3410-3510;
Fax, 82-2-3410-4400; E-mail, jeholee@gmail.com
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Abbreviations: AGR2, anterior gradient 2; *agt*, angiotensinogen; RT-CES, real-time cell electronic sensing system; *THBS1*, thrombospondin 1; *vegf A*, vascular endothelial growth factor A

Abstract

Ovarian cancer is a leading cause of death in women. Early detection of ovarian cancer is essential to decrease mortality. However, the early diagnosis of ovarian cancer is difficult due to a lack of clinical symptoms and suitable molecular diagnostic markers. Thus, identification of meaningful tumor biomarkers with potential clinical application is clearly needed. To search for a biomarker for the early detection of ovarian cancer, we identified human anterior gradient 2 (AGR2) from our systematic analysis of paired normal and ovarian tumor tissue cDNA microarray. We noted a marked overexpression of AGR2 mRNA and protein in early stage mucinous ovarian tumors compared to normal ovarian tissues and serous type ovarian tumors by Western blot analysis and immunohistochemistry. To further elucidate the role of AGR2 in ovarian tumori-

genesis, stable 2774 human ovarian cancer cell lines overexpressing AGR2 were established. Forced expression of AGR2 in 2774 cells enhanced the growth and migration of ovarian cancer cells. AGR2 protein was detected in the serum of mucinous ovarian cancer patients by Western blot and ELISA analysis. Thus, AGR2 is a potential biomarker for the diagnosis of mucinous ovarian cancer and an ELISA assay may facilitate the early detection of mucinous ovarian cancer using patient serum.

Keywords: AGR2 protein, human; biological markers; ovarian neoplasms

Introduction

Ovarian cancer is one of the most lethal gynecologic malignancies and has among the worst of prognoses. The lethality of ovarian cancer is primarily attributable to the advanced stage of the disease at the time of the initial diagnosis. Although the cure rate from currently available therapies is as high as 90% when the cancer is detected in an early stage (American Cancer Society, 2005), the majority of early-stage cancers are asymptomatic and most cases are diagnosed at an advanced stage. Thus, long-term survival can be achieved in < 30% of patients (Edwards *et al.*, 2005). Therefore, screening for an early-stage diagnosis can significantly reduce mortality. The identification of reliable diagnostic biomarkers for the early detection of ovarian cancer is critical for reducing the mortality rate of ovarian cancer. Furthermore, identification of meaningful tumor biomarkers with potential clinical application would provide useful information regarding patient diagnosis, prognosis, and possible therapeutic options.

Anterior gradient homolog 2 (AGR2), also known as hAG-2 (Fletcher *et al.*, 2003) and Gob-4 (Komiya *et al.*, 1999), is the human orthologue of the secreted *Xenopus laevis* anterior gradient protein, XAG-2. XAG-2 has a putative role in ectodermal patterning of the frog embryo and is regulated by *noggin* and *chordin* (Aberger *et al.*, 1998). The expression of XAG-2 induces cement gland differentiation and expression of neural marker genes in a fibroblast growth factor-dependent manner (Aberger *et al.*, 1998). Up-regulation of the AGR2 gene has been reported in breast cancer (Fri-

tzsche *et al.*, 2006), prostate cancer (Kristiansen *et al.*, 2005; Zhang *et al.*, 2005), and endometrial and renal carcinomas (Fritzsche *et al.*, 2006). AGR2 has been proposed as one of the potential markers to detect circulating tumor cells in the peripheral blood in patients with metastatic cancers, including colorectal, prostate, and breast cancers (Smirnov *et al.*, 2005). The expression of AGR2 has also been described to be associated with a positive estrogen receptor status of tumor cells. However, the expression and role of AGR2 in mucinous ovarian cancer remains uncharacterized and requires further study.

Initially, we set out to identify potential biomarkers that could be used as a non-invasive method of early diagnosis of ovarian cancer by cDNA microarray analysis. Herein we report that the secreted protein, AGR2, is highly overexpressed in human mucinous ovarian tumor tissues and patient sera compared with tissue and sera samples from healthy women, as determined by biochemical and molecular approaches. To further explore the roles of AGR2 in ovarian tumorigenesis, stable AGR2-overexpressing human ovarian cancer cells were established and characterized. The findings from this study suggest that AGR2 can serve as a potential diagnostic marker for the detection of mucinous ovarian cancer and AGR2 detection may provide an efficient and sensitive tool for the early detection of human mucinous ovarian cancer.

Results

Identification of a novel biomarker for human mucinous ovarian cancer

To identify novel molecular markers that are more sensitive and specific for the early detection of ovarian cancer, cDNA microarray was carried out and analyzed with 32 tissue samples from ovarian cancer patients. Among the up-regulated genes, AGR2 was chosen because the subcellular localization of AGR2 using the PSORT II program revealed that it is a secretory protein. AGR2 was markedly up-regulated in all mucinous-type ovarian cancers, but no significant expression (< 25% of all cases) was observed in serous-type ovarian cancers compared with normal ovarian tissues (Figure 1). A scatter plot shows a > 5-fold up-regulation of AGR2 mRNA in mucinous ovarian tumors, whereas the majority of serous-type ovarian tumors had a < 5-fold up-regulation (Figure 1A). A further comparison between adenocarcinoma and borderline tumors in each tumor type clearly revealed preferential expression of

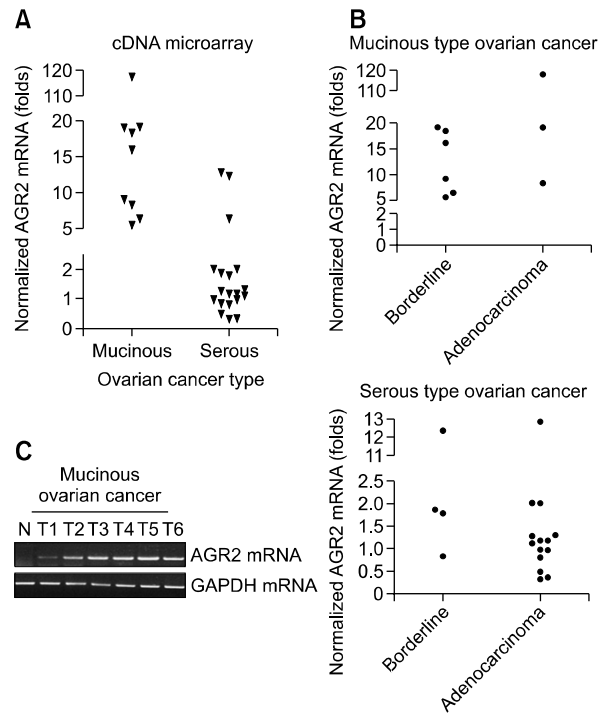


Figure 1. Up-regulation of AGR2 mRNA in human mucinous ovarian cancer. (A, B) Scatter plot analysis of cDNA microarray. AGR2 expression was analyzed according to the tumor type (A) and cancer grade (B). Scattered plot shows > 5-fold up-regulation of AGR2 mRNA in mucinous ovarian tumors, whereas the majority of serous-type ovarian tumors had a < 5-fold up-regulation. (C) RT-PCR analysis of AGR2 expression in mucinous ovarian cancer tissues. Total RNA was used to confirm the up-regulation of AGR2 by cDNA microarray in mucinous ovarian tumor tissues. GAPDH mRNA was used as a control to normalize the level of mRNA in each sample. N: normal ovary.

AGR2 in mucinous ovarian tumors compared with serous ovarian tumors (Figure 1B). Next, to confirm the expression of AGR2 mRNA in mucinous ovarian cancers, RT-PCR analysis was performed in six different human mucinous ovarian cancer tissues and one normal human ovarian tissue (Figure 1C). In agreement with microarray data, human mucinous cancer had a markedly increased expression of AGR2 mRNA.

To determine the level of expression of AGR2 protein, 20 ovarian tumors and 4 normal healthy ovarian tissues were immunostained with anti-AGR2 antibody (Figure 2A). AGR2 protein was detected in the cytoplasm of all tissues. AGR2 protein was expressed at a basal level in normal ovary surface epithelium (lane 1) and serous ovarian tumor tissues (lanes 4-5), whereas a highly elevated level of AGR2 protein expression was detected in mucinous ovarian tumors (lanes 2-3; Figure 2B). These results suggest that AGR2 protein may be a good marker for the detection of mucinous ovarian tumors. AGR2 protein ex-

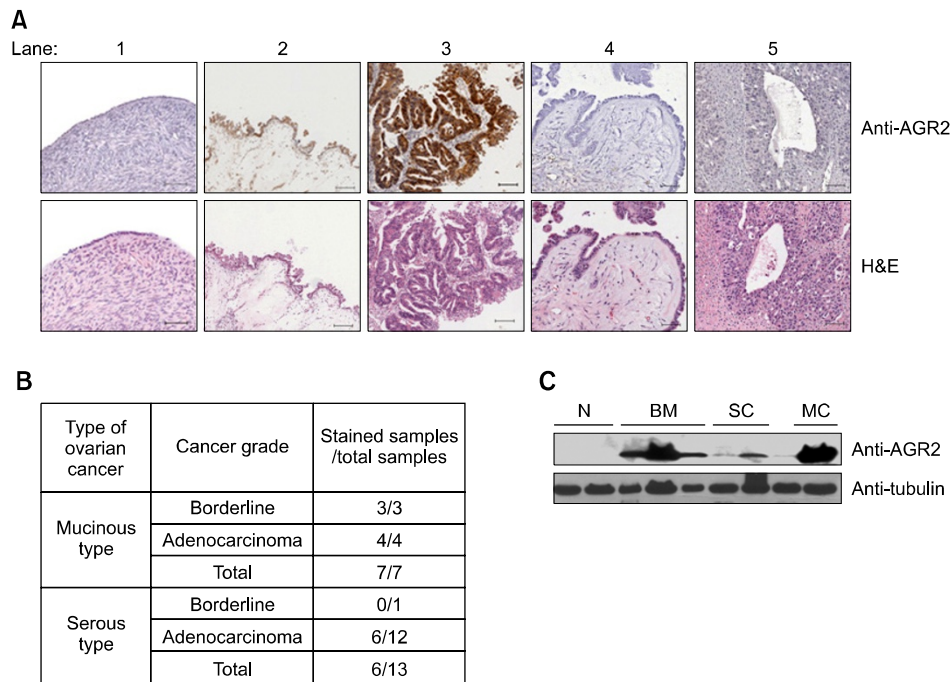


Figure 2. Overexpression of AGR2 protein in human mucinous ovarian cancer. (A) Top panels: Immunohistochemistry of AGR2 protein in various types of ovarian tumor tissues. The sections were counter-stained with hematoxylin as follows: 1) normal ovary; 2) borderline mucinous tumor of the ovary; 3) mucinous adenocarcinoma of the ovary; 4) borderline serous tumor of the ovary; and 5) serous adenocarcinoma of the ovary. Bottom panels: Corresponding H&E staining of respective tumor section (Scale bar: 50 μ m). (B) Summary of AGR2 protein expression in ovarian cancer by immunohistochemistry. AGR2 expression was further analyzed according to the tumor type and cancer grade. (C) Expression of AGR2 protein in ovarian tumor by Western blot analysis. Total tissue lysates were prepared in RIPA buffer and then subjected to Western blot analysis with anti-AGR2 antibody. N, Normal ovary; BM, Borderline mucinous tumor of the ovary; SC, Serous adenocarcinoma of the ovary; MC, Mucinous adenocarcinoma of the ovary.

pression in ovarian tumor tissues was further confirmed by Western blot analysis (Figure 2C). The anti-AGR2 antibody detected a predominant 18 kDa protein in the protein extracts from mucinous ovarian tumors. In contrast, there was weak or no detectable AGR2 protein in normal ovarian tissues. These results are consistent with immunohistochemical staining of AGR2 protein, and very high expression of AGR2 protein in mucinous ovarian tumors compared with no staining in normal counterparts. These results confirmed the markedly elevated AGR2 expression of protein in mucinous ovarian tumor tissues.

Generation of human ovarian cancer cells overexpressing human AGR2

To further elucidate the role and mechanism of AGR2 in ovarian tumorigenesis, human AGR2-overexpressing ovarian cancer cells were generated by G418 selection (Figure 3A). Control vector- and AGR2-expressing stable cells were tested for the expression of AGR2 protein by

Western blot analysis and confocal microscopy after indirect immunofluorescence staining (Figure 3B). Compared to control vector stable transfectants, AGR2 stable transfectants (clones A1 and A2) exhibited expression of AGR2 protein. AGR2 stable transfectants showed strong cytoplasmic staining of AGR2 protein by confocal imaging analysis, whereas control vector stable cells had no staining for AGR2 protein (Figure 3B). The subcellular localization of AGR2 protein was predicted to be cytoplasmic and secreted using the PSORT II program (<http://psort.nibb.ac.jp>). To clarify the subcellular localization of overexpressed AGR2 protein, 2774-AGR2 stable transfectants were biochemically fractionated, then AGR2 protein expression was assessed by Western blot analysis with anti-AGR2 antibody (Figure 3C). AGR2 protein was preferentially expressed in the cytoplasm. Furthermore, to confirm secretion of AGR2 protein in the serum-free conditioned medium, conditioned medium from the AGR2 stable transfectants were analyzed by Western blot analysis with anti-AGR2 antibody (Figure 3D). As

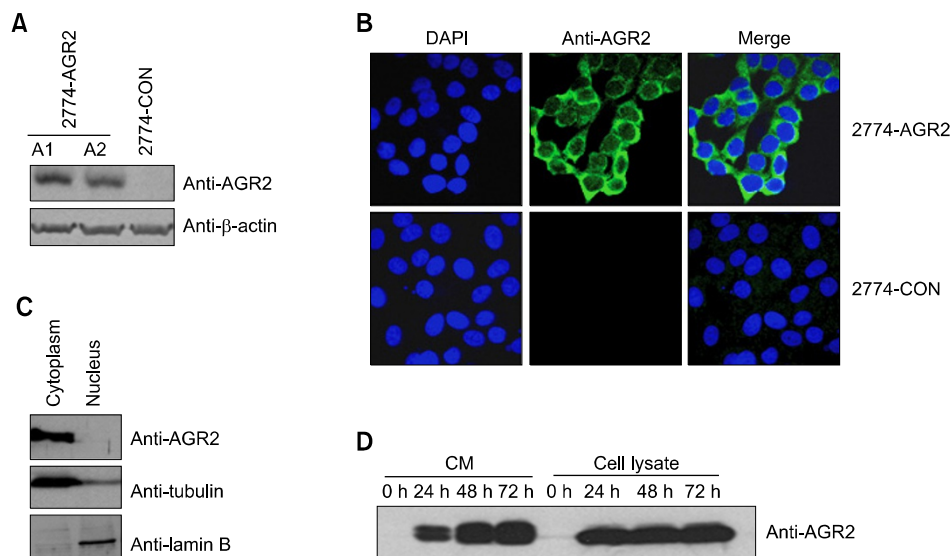


Figure 3. Establishment of AGR2 overexpressing human 2774 ovarian cancer stable transfectants. (A) Expression of AGR2 protein in 2774 stable transfectants. Human AGR2 mammalian expression vector was transfected into 2774 cells, then AGR2-overexpressing stable cells were selected by G418 selection. Stable 2774 transfectants were assessed for AGR2 protein expression by Western blot analysis with anti-AGR2 antibody. (B) Subcellular localization of AGR2 in AGR2 stable transfectants. Vector control and AGR2 stable transfectants were processed for indirect immunofluorescence microscopy for subcellular localization of AGR2 protein in human 2774 ovarian cancer cells. AGR2 protein was expressed in all AGR2 stable transfectants and showed a strong preferential cytoplasmic localization. (C) Subcellular localization of AGR2 protein. Whole cell lysates prepared in RIPA buffer from the stable 2774 cells were fractionated, then subjected to Western blot analysis with anti-AGR2 antibody. AGR2 protein was detected in the cytoplasmic fraction. Subcellular fractionation was confirmed by Western blot analysis with anti-lamin B and anti-tubulin antibodies. (D) AGR2 protein is a secreted protein. Conditioned medium was collected from the indicated stable AGR2 transfected 2774 cells at the indicated incubation time, then analyzed by Western blot analysis with anti-AGR2 antibody.

expected, we were able to detect AGR2 protein in the serum-free conditioned medium in AGR2 stable transfectants.

Enhancement of ovarian cancer cell proliferation and migration by AGR2 overexpression

To determine the effect of AGR2 overexpression in human ovarian cancer cells, AGR2-expressing stable 2774 ovarian cancer cells were tested for the effect on cancer cell growth by a real-time cell electronic sensor system (CES). Forced expression of AGR2 increased ovarian cancer cell growth and proliferation (Figure 4A). Next, the effect of AGR2 on cancer cell migration was assessed using a Boyden chamber. Overexpression of AGR2 significantly increased cancer cell migration that is necessary for cancer cell invasion (Figure 4B).

To gain insight into the function of AGR2 in ovarian tumorigenesis, total RNA extracted from control vector and AGR2 stable cells was used for cDNA microarray analysis to identify the target genes for further functional studies. cDNA microarray analysis of AGR2 stable transfectants revealed that AGR2 overexpression up-regulated

the expression of genes involved in cell proliferation, invasion, and angiogenesis, which is important for tumor progression and metastasis. For example, expression of vascular endothelial growth factor A (*vegf A*) and angiotensinogen (*agt*) genes, which are secreted and involved in angiogenesis, were elevated 3.44- and 4.45-fold, respectively, whereas interleukin-8 (*IL-8*) and thrombospondin 1 (*THBS1*), which are involved in cell adhesion and negative regulation of cell proliferation, were each reduced 2-fold (Table 1). Furthermore, the genes involved in cell death and cell cycle regulation were down-regulated. The genes involved in cell adhesion, such as *P-cadherin* and *THBS1*, were down-regulated.

Detection of AGR2 protein in the serum from ovarian cancer patients

Serum samples were analyzed for the detection of AGR2 protein by Western blot analysis with anti-AGR2 antibody (Figure 5A). Equal protein loading of serum samples was confirmed by Ponceau S staining of the same blot (bottom panel). AGR2 protein expression was significantly elevated in the

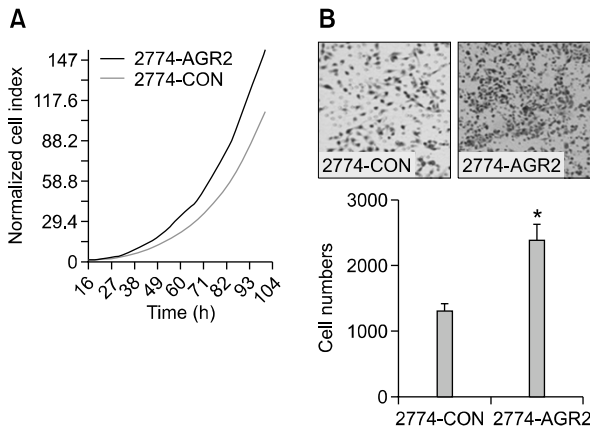


Figure 4. AGR2 overexpression increased cell growth and cell migration of human ovarian cancer cells. (A) Control vector- and AGR2-over-expressing stable 2774 cells were plated onto 96-well sensor plates at 2.5×10^3 cells per well, then incubated overnight. Growth of cells was continuously monitored using real-time cell electronic sensing (RT-CES) system. (B) Enhancement of cancer cell migration by AGR2 over-expression. Cancer cell migration was assayed using Transwells after coating the lower surface of the filter with gelatin. The number of cancer cells that migrated to the lower side of the filter was counted. The results are the means \pm SD (*, $P < 0.05$ compared to 2774-CON).

serum of the mucinous-type ovarian cancer, as shown by Western blot analysis (top panel). Expression of AGR2 protein in patient serum was also analyzed by ELISA methods. Five mucinous-type ovarian cancer serum samples, 9 serous-type samples, and 11 normal healthy women samples were assessed for the detection of AGR2 by ELISA and summarized (Figure 5B). The elevated expression of serum AGR2 protein was detected in

mucinous ovarian tumor sera compared with normal and serous-type ovarian tumor sera by ELISA. The significance of the results was assessed by the Kruskal-Wallis rank-sum test ($P < 0.0034$).

Discussion

Cancer cells are known to acquire the hallmarks of malignancy through the accumulation of advantageous gene activation and inactivation events over long periods of time (Hanahan and Weinberg, 2000). To identify a novel ovarian tumor marker for early detection, pooled scrapings of human normal ovarian surface epithelium rather than whole ovaries were used to compare gene expression with ovarian tumors. Previously, AGR2 was reported to be expressed in cancers of the breast (Fritzsche *et al.*, 2006), pancreas (Ramachandran *et al.*, 2008; Riener *et al.*, 2009), and prostate (Zhang *et al.*, 2005). Overexpression or suppression of AGR2 was shown to affect cell proliferation, cell invasion, metastasis, and tumor growth of pancreatic cancer (Ramachandran *et al.*, 2008). Recent reports support a role for AGR2 in cellular transformation and growth of esophageal adenocarcinoma (Wang *et al.*, 2008).

Our systematic approach combining cDNA microarray analysis, RT-PCR, immunohistochemistry, Western blot analyses, and bioinformatics facilitated the identification of AGR2 as a novel candidate of an ovarian tumor marker for the early detection of mucinous ovarian tumors. More importantly, in terms of clinical usefulness, we were

Table 1. cDNA microarray analysis of AGR2 stable cells

A. Genes up-regulated in AGR2 stable cells						
GenBank	Gene	Gene description	Normalized ratio	Subcellular localization	Biological function	
NM_002961	<i>S100A4</i>	S100 calcium binding protein A4	2.43	Cytoplasm	Invasive growth	
NM_005980	<i>S100P</i>	S100 calcium binding protein P	5.01	Cytoplasm	Cell life, proliferation/growth	
NM_003225	<i>TFF1</i>	Trefoil factor 1	4.19	Secreted	Cell growth and/or maintenance	
NM_001007139	<i>IGF2</i>	Insulin-like growth factor 2	3.84	Secreted	Cell growth	
NM_001025366	<i>VEGFA</i>	Vascular endothelial growth factor A	3.44	Secreted	Angiogenesis	
NM_000029	<i>AGT</i>	Angiotensinogen	4.45	Secreted	Angiogenesis	
B. Genes down-regulated in AGR2 stable cells						
GenBank	Gene	Gene description	Normalized ratio	Subcellular localization	Biological function	
NM_001717	<i>BNC1</i>	Basonuclin 1	0.25	Nucleus	Regulation of transcription, DNA dependent	
BC016333	<i>CCNL2</i>	Cyclin L2	0.47	Nucleus	Regulation of cell cycle	
NM_001793	<i>CDH3</i>	Cadherin 3, type 1, P-cadherin	0.29	Integral to membrane	Cell adhesion	
NM_001425	<i>EMP3</i>	Epithelial membrane protein 3	0.39	Integral to membrane	Cell death	
NM_003246	<i>THBS1</i>	Thrombospondin 1	0.40	Extracellular matrix	Cell adhesion	
NM_000584	<i>IL8</i>	Interleukin 8	0.50	Extracellular space	Negative regulation of cell proliferation	

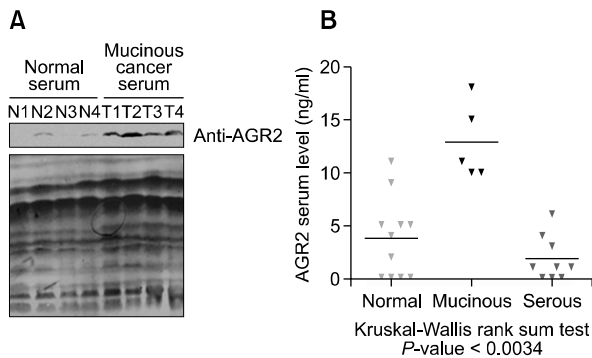


Figure 5. Detection of AGR2 protein in the serum from mucinous ovarian cancer patients. (A) Serum AGR2 level was assessed by Western blot analysis of de-albuminized sera derived from the normal volunteers and tumor patients after processing using immunoaffinity albumin and IgG depletion kit. Equal serum protein loading was confirmed by staining the same blot with Ponceau S. (B) Twenty-five sera (5 mucinous-type ovarian cancer, 9 serous-type ovarian cancer, and 11 normal ovarian samples) were processed for the detection of AGR2 protein by ELISA assay and summarized by scatter plot analysis. Statistical significance was analyzed by the Kruskal-Wallis rank sum test ($P < 0.005$).

able to detect AGR2 in patient serum by Western blot and ELISA assays. Detection in patient serum by an ELISA assay developed in this study provides an efficient screening of mucinous ovarian tumors among other ovarian tumors. Cumulative molecular genetic analyses of ovarian cancers have revealed genetic alterations of several genes in ovarian tumors, such as c-erb-B2, c-myc, and p53 (Aunoble *et al.*, 2000). However, none of these mutations are diagnostic of malignancy. Previous reports by other investigators have identified multiple molecular targets that differ in normal and malignant cells in epithelial ovarian cancer, including cell cycle regulators, growth factor receptors, signal transduction pathways, genes that regulate drug resistance, and angiogenesis (reviewed in See *et al.*, 2003; Sharma and Odunsi, 2005). In addition, AGR2 was shown to be a useful marker for detection of circulating tumor cells (CTCs) in peripheral blood of advanced cancer patients (Smirnov *et al.*, 2005). The AGR2 protein has been previously associated with breast carcinomas by suppression subtractive cDNA library screening between a malignant human breast cancer cell line (MCF-7) and a benign breast tumor-derived cell line (Huma 123; Liu *et al.*, 2005). Screening of breast tumor specimens using RT-PCR and immunocytochemistry with anti-AGR2 antibodies has demonstrated that the presence of AGR2 mRNA and protein is statistically and significantly associated with ER α -positive carcinomas and malignancy (Fritzsche *et al.*, 2006). In addition, expression of AGR2 has been reported to be

strongly associated with markers of tumor differentiation (estrogen receptor positivity and lower tumor grade). Although CA-125 is the best clinical marker for ovarian cancer in the postmenopausal age group, CA-125 has proven to be a poor diagnostic tumor biomarker for early stage ovarian cancer (Liede *et al.*, 2002). Importantly, the CA-125 level was not elevated in the mucinous subtype. Previous efforts to characterize serum biomarkers for the detection of early-stage ovarian cancer identified transthyretin, beta-hemoglobin, apolipoprotein A1, and transferrin using micro-LC-MS/MS (Kozak *et al.*, 2005). Unfortunately, the combination of CA-125 with biomarkers, such as transthyretin, beta-hemoglobin, apolipoprotein A1, and transferrin, did not further improve the detection of mucinous tumors (Kozak *et al.*, 2005). There has been an increase in the search for ovarian cancer markers using SELDI-TOF-MS with the use of multiple markers (Zhang *et al.*, 2004), but there is still no significant improvement for detection of mucinous ovarian cancer. Considering the preferential expression of borderline mucinous cancer, AGR2 would be beneficial to improve the early detection of mucinous ovarian tumors.

To gain insight into the role of AGR2 in ovarian tumorigenesis, AGR2-overexpressing stable human ovarian cancer cells were generated and our results showed that AGR2 enhanced cell growth and migration of ovarian cancer cells. Consistent with the involvement of AGR2 in cell growth and migration, our cDNA microarray analysis of AGR2-stable cells revealed that AGR2 overexpression up-regulated the expression of genes involved in cell proliferation, invasion, and angiogenesis, which is important for the tumor progression and metastasis characteristic of ovarian cancer. By contrast, the genes involved in the negative regulation of cell proliferation, cell adhesion, and cell death were down-regulated. Moreover, AGR2 protein can be detected in the serum of mucinous ovarian tumor patients by Western blot and ELISA methods. These results suggest that serum AGR2 can be a biomarker for diagnosis and prediction of mucinous ovarian cancers. This is the first report that AGR2 is a candidate molecular marker for mucinous ovarian tumors. However, validation of AGR2 as a potential serum marker is needed with a large-scale analysis of ovarian cancer blood samples and can be compared with other validated serum markers. We continue to expand our effort to collect serum from mucinous ovarian cancer patients and normal volunteers. Because AGR2 has previously been reported to be associated with tumor metastasis, it would be interesting to determine the role of AGR2 in ovarian tumor pro-

gression and metastasis. Analysis of AGR2 protein suggests potential protein modification that might be important for AGR2 function (Park and Lee, unpublished data). Further functional analysis of AGR2 protein modification during tumor progression would provide information for its functional significance in addition to its role in diagnosis. Recently AGR2 was classified as a member of the protein disulfide isomerase (PDI) family and shown to play a special role in the production of mucus (Park *et al.*, 2009). AGR2 is also expressed in mucus-producing cells, and AGR2-deficient mice were unable to produce intestinal mucin and became highly susceptible to experimentally-induced colitis.

The impact of results in this study on cancer biomarker discovery efforts is significant because ovarian cancer remains a highly fatal cancer due to the lack of reliable screening tests and high recurrence of ovarian cancer. The results from this study suggest that AGR2 can be a potential biomarker for diagnosis of mucinous ovarian cancer and the ELISA assay would facilitate the early detection of mucinous ovarian cancer using a non-invasive method using patient serum.

Methods

Cell culture

Human ovarian cancer (2774 cells) and human fetus kidney epithelial cells (HEK 293 cells) were cultured in DMEM and EMEM, respectively, and supplemented with 10% FBS and antibiotics (penicillin [100 units/ml] and streptomycin [100 µg/ml]; Invitrogen Corporation, Carlsbad, CA) in a 5% CO₂ humidified atmosphere at 37°C.

Tissue and serum protein preparation

All human tissue and serum samples were collected with institutional approval at Samsung Medical Center. All ovarian tissues, including normal and tumor sources, were collected from women who underwent surgery in the Department of Obstetrics and Gynecology at Samsung Medical Center. The tissue samples were immediately stored in liquid nitrogen and stored at -80°C until analysis. Paraffin archives from the Department of Pathology of Samsung Medical Center were used to confirm the expression of AGR2 by immunohistochemistry. All serum samples were collected by venipuncture from patients and healthy donors. Sera from women with ovarian cancer were collected before surgical removal of the ovaries. Both the stage and histology of the cancer were determined by standard pathologic methods. Serum was separated, centrifuged (3000 × g, 20 min), and stored at -80°C in 50 µl aliquots until use. Serum specimens were used to detect the serum AGR2 protein level by Western blot analysis after depleting albumin and IgG from serum samples using a Vivapure anti-HAS/IgG kit (Sartorius Stedim Biotech,

Bohemia, NY) according to the manufacturer's protocol.

Chemicals

Antibodies against AGR2 were purchased from Abnova (clone 1C3; Taipei city, Taiwan) and Abcam (rabbit polyclonal; Cambridge, UK). Tubulin and lamin B were purchased from Santa Cruz Biotechnology, Inc. (Santa Cruz, CA), and β-actin from Sigma (St. Louis, MO). 4',6-diamidino-2-phenylindole (DAPI) was purchased from Roche Applied Science (Mannheim, Germany). All other chemical reagents were purchased from Sigma, unless otherwise indicated.

Stable transfection and generation of AGR2-overexpressing human ovarian cells

Human 2774 ovarian cancer cells were transiently transfected with control vector or AGR2-expressing vector using Effectene (Qiagen, Hilden, Germany) according to the manufacturer's instructions. In brief, 2774 cells were seeded the day before transfection, then incubated until reaching 60% confluence on the day of transfection. After selection in medium containing G418 (500 µg/mL; Invitrogen), G418-resistant discrete colonies were selected for further expansion in selection medium. Several clones for vector control and AGR2-overexpressing 2774 cells were maintained with DMEM containing G418.

RNA isolation, cDNA microarray analysis, and RT-PCR

Total cellular RNA was extracted from ovarian tumor and matched normal tissues obtained from 32 patients using TRIZOL reagent (Invitrogen Life Technologies), according to the manufacturer's protocol. Human 17K cDNA microarrays (GenomicTree, Daejeon, Korea) containing 17,075 human cDNAs representing 15,724 different genes (UniGene clusters) were used in gene expression analysis, as previously described (Yang *et al.*, 2003). Briefly, each 50 µg of total RNA was mixed with 2 µg of oligo-(dT)₂₄ (GenoTech, Daejeon, Korea) and incubated at 65°C for 10 min. Single-stranded cDNA was synthesized in the presence of Cy3-dUTP or Cy5-dUTP (1 mM each, NEN Life Science Products, Boston, MA) at 42°C for 2 h. Total RNAs from ovarian tumors and normal ovary tissues were labeled with Cy5 and Cy3, respectively. Both Cy5- and Cy3-labeled cDNA were purified using PCR purification kit (Qiagen) and hybridized onto human 17K cDNA microarrays. The arrays were incubated at 42°C for 16 h in the humidified hybridization chamber. The hybridized microarrays were imaged by using the Axon 4000B scanner (Molecular Devices, CA). The signals and background fluorescence intensities were calculated for each probe by averaging the intensities of every pixel inside probe spot region using GenePix Pro 4.0 software (Molecular Devices). Spots of poor quality or indistinguishable signal levels from the background were excluded. Data normalization and selection of differentially expressed genes were performed using GeneSpring 7.3.1 (Agilent Technologies, Palo Alto, CA). To analyze differentially expressed

genes between tumor and paired normal ovary tissues, the "Cross gene error model for replicates" was used. Gene expression data was normalized using global LOWESS normalization. Probes which were changed ≥ 2.0 -fold ratio between tumor and control normal ovary tissues were selected and considered as differentially-expressed probes. For RT-PCR analysis, 1 μ g of total RNA was dissolved in distilled water, then used for cDNA synthesis by superscript reverse transcriptase (RT; Invitrogen), then PCR amplification was carried out with specific AGR2 primers as follows: 94°C for 3 min, followed by 20 cycles at 94°C for 30 s, 55°C for 30 s, and 72°C for 1.5 min. The following primers were used for PCR amplification: forward primer, 5'-GGGATGGAGAAAATTCCAGTG-3'; and reverse primer, 5'-GGGTACAATTCAGTCTTCAG-3'. PCR products were separated by 1.2% agarose gel electrophoresis. GAPDH was used as a loading control.

Subcellular fractionation

Subcellular fractionation was carried out using a ProteoExtract Subcellular Proteome Extraction kit (Calbiochem, Darmstadt, Germany). Cultured cells were harvested with 500 μ l of cell lysis buffer (10 mM Tris-HCl [pH 7.5], 10 mM NaCl, 1 mM KH₂PO₄, 5 mM NaHCO₃, 1 mM CaCl₂, and 0.5 mM MgCl₂) with a protease inhibitor cocktail (Roche Diagnostics Corp. Penzberg, Germany). Cells were allowed to swell for 5 min, then homogenized 50 times using a Dounce homogenizer (Wheaton, Millville, IL). Lysates were centrifuged for 10 min at 7,500 rpm in a centrifuge at 4°C, and the cytoplasmic fraction was transferred to a clean tube. The nuclear pellet was washed once with cell lysis buffer before resuspending with SDS-loading buffer. The efficiency of subcellular fractionation was confirmed by Western blot analysis with anti-lamin B and anti-tubulin as nuclear and cytoplasmic protein marker antibodies, respectively.

Western blot analysis

Cultured cells and tissue lysates were lysed by sonication in phosphate-buffered saline with RIPA buffer (50 mM Tris-HCl [pH 7.5], 150 mM NaCl, 1% NP40, 0.5% sodium deoxycholate, and 0.1% SDS) containing protease inhibitors (protease inhibitor mixture; Roche Applied Science), as described previously (Park *et al.*, 2005). Proteins were separated by SDS-PAGE and electrophoretically transferred onto an ECL nitrocellulose membrane (GE Healthcare, London, UK). Western blot analysis was performed with the primary antibody (1:1000 or 1:2000 dilution) and an enhanced chemiluminescence system (ECL; Amersham Sciences, Chicago, IL).

Indirect immunofluorescence staining and confocal microscopy

Cells were grown on sterilized glass coverslips, fixed with 4% paraformaldehyde in 0.1 M phosphate buffer for 15 min, then blocked with 0.1% BSA in PBS. Then cells were incubated with anti-AGR2 (1:500 dilution) in PBS, followed by incubation with Alexa 488-conjugated secondary antibody (1:5000 dilution; Vector, Burlingame, CA). The slides

were washed three times with PBS, and mounted in mounting media (Vectorshield; Vector). Images were captured with a Bio-Rad confocal microscope (Bio-Rad Laboratories, Hercules, CA).

Immunohistochemistry

Immunohistochemical analysis was carried out on a formalin-fixed, paraffin-embedded specimen cut at a thickness of 4 μ m. The paraffin block specimens used in the current experiments were derived from archived tissue (not representing samples from patients who contributed fresh specimens) derived from patients having surgery at the Samsung Medical Center. All slides were deparaffinized in xylene, rehydrated through a graded alcohol series, and washed in PBS. The slides were treated with 0.3% hydrogen peroxide in PBS for 5 min at room temperature to quench the endogenous peroxidase activity. For antigen retrieval, sections were microwaved twice for 10 min in 10 mM citrate buffer (pH 6.0). For AGR2 staining, polyclonal rabbit antisera against peptides derived from the AGR2 protein sequence (dilution 1:250; Abcam) were used and the primary antibody was incubated at 4°C overnight. As a negative control, slides were also processed without primary antibody. Detection was done by the conventional streptavidin-biotin method with 3,3'-diaminobenzidine tetrahydrochloride as a chromogen (DAKO, Copenhagen, Denmark). The slides were counterstained with hematoxylin for 5 min, then dehydrated by sequentially washing in an ethanol series and rinsing three times with xylene. The slides were then mounted with Consul-mount (Thermo Scientific, Waltham, MA).

Subcloning of human AGR2

The full-length coding sequence of human AGR2 was PCR-amplified using AGR2 primers, as described below. Thermal cycling conditions comprised an initial step at 95°C for 5 min, followed by 20 cycles at 95°C for 1 min, 55°C for 1 min, 72°C for 2 min, then a final extension at 72°C. The forward primer was 5'-GGGGAATTCATGGAGAAAATTCCAGTG-3' and the reverse primer was 5'-GGGCTCGAGTTACAATTCAGTCTTCAG-3'. Amplified full-length AGR2 was then subcloned into EcoR1-Xho I-digested mammalian expression vector pcDNA3.1 Myc-His (Invitrogen). The sequence of AGR2 in mammalian vector was confirmed by DNA sequence analysis.

Real-time cell growth monitoring

The effect of AGR2 overexpression on cell growth was assessed using the real time electronic sensing system (RT-CES; ACEA Biosciences, San Diego, CA), according to the manufacturer's protocol. Control vector- and AGR2-overexpressing stable 2774 cells were plated onto 96-well sensor plates at 2.5×10^3 cells per well, then incubated overnight. Cells were continuously monitored using a RT-CES.

In vitro migration assays

Migration of ovarian cancer cells was assessed using a

Biocoat matrigel chamber (Becton Dickinson Labware, Bedford, MA), as previously described (Choi *et al.*, 2005). Briefly, 1×10^5 cells were plated in each well and the bottom chamber outside the transwell was filled with 500 μ l of DMEM medium supplemented with 1% BSA. After incubation for 16 h, the cells were fixed in 100% methanol for 5 min and stained in 0.1% eosin. The number of cells that migrated to the lower side of the filter was counted.

Sandwich ELISA assay

To develop a sensitive sandwich ELISA assay for AGR2 protein detection in patient serum, optimized dilutions of monoclonal AGR2 antibodies were coated onto 96-well Immobilon plates overnight at 4°C in $1 \times$ PBS buffer. After washing, plates were blocked with 0.05% Tween 20/2.5% BSA/1 \times PBS at RT for 1 h. Patient serum was added to the well, then incubated for 2 h at RT. After washing unbound antigen solutions, the second anti-AGR2 antibody was added and incubated for 4 h. After washing unbound antibody, HRP-conjugated goat anti-mouse IgG was added at a 1:5000 dilution. Detection was performed with Super AquaBlue ELISA substrate (eBioscience, San Diego, CA) and stopped with 0.5 M sulfuric acid. Plates were read in a Tunable microplate reader (Molecular Devices) at 405 nm and analysis was performed using the SoftMax Pro v4.3 LS software (Molecular Devices).

Statistical analysis

Experiments were independently performed three or more times. The data are presented as the mean \pm standard deviation. An unpaired Student's *t*-test was used to compare the means of cell migration between the vector control and AGR2-overexpressing 2774 stable cells. The Kruskal-Wallis rank-sum test was used to evaluate the significant difference in AGR2 serum levels between normal volunteers and ovarian cancer patients. *P*-values < 0.05 were considered statistically significant.

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