Abstract. In light of the poor prognosis for ovarian cancer, research continues for innovative and efficacious treatment modalities. It is now widely accepted that new approaches for the treatment of ovarian cancers are pivotal to further improve the prognosis of this disease. We investigated the role of HAI-1 and HAI-2, and evaluated their clinical importance in ovarian cancer. The distribution of cases that scored positive for each of the biological parameters examined was correlated with HAI-1 and HAI-2 expression status obtained by immunohistochemistry. There was a significant correlation between HAI-2 expression and stage (p=0.031), amount of ascites (p=0.002) and diameter of residual tumor (p=0.034). HAI-1 expression was a significant associated with stage (p=0.040). Furthermore, the low HAI-1 and HAI-2 expression was a significant predictor for poor prognosis when compared with high HAI-1 and HAI-2 expression (disease-free survival rate; p=0.031 and p=0.003, overall survival rate; p=0.048 and p=0.001). Therefore, we investigated biological functions and effects of HAI-1 and HAI-2 using OVCAR-3 ovarian cancer cell lines. HAI-1 and HAI-2 show potential inhibitory effects mediated by reduction of matriptase and hepsin expression which leads to apoptosis through increasing the level of Bak and reducing the level of Bcl-2 on ovarian cancer. These findings indicate that low HAI-1 and HAI-2 expression in ovarian cancer may be associated with poor prognosis. HAI-1 and HAI-2 could be considered a therapeutic target for treatment approaches in ovarian cancer.

Introduction

Despite its pervasive impact, the etiology of ovarian cancer and the factors that promote its progression are not yet well understood. It is now widely accepted that new approaches for the treatment of ovarian cancers are pivotal in further improving prognosis of this disease.

Hepatocyte growth factor (HGF) is secreted from mesenchymal cells of the liver as an inactive single-chain form and normally remains in this form associated with the extracellular matrix (1). It has mitogenic, motogenic, morphogenic functions in various types of cells through its high-affinity receptor tyrosine kinase that is encoded by the c-Met proto-oncogene. The c-Met have important roles in migration, invasion of tumor cells, differentiation, and tumor angiogenesis (2,3). The c-Met overexpression in ovarian cancer is associated with ovarian cancer progression, and high c-Met expression correlates with poor survival (4,5).

HGF is secreted as an inactive pro-peptide which is cleaved by HGF activator (HGFA) into its active form. HGFA is regulated by two inhibitors, HGFA inhibitor type I (HAI-1) and type II (HAI-2) they are type I transmembrane proteins with two Kunitz-type serine protease domains in the extracellular portion, and a cell surface expression on epithelial cells (6-9). Among HAI-1 and HAI-2, membrane-form HAI-1 can bind to HGFA on the cellular surface, but membrane-form HAI-2 cannot (10). It is thus very likely that HAI-1 and HAI-2 have distinct biological roles in vivo.

The binding of HGFA to HAI-1 is reversible and may reserve available HGFA on the cell surface to ensure concentrated pericellular HGFA activity (11). HAI-1 appears to be the cognate inhibitor of HGFA and matriptase, and both HGFA and hepsin (11,12). HAI-2 is a more efficient inhibitor of hepsin and displays a broader inhibitory spectrum than HAI-1 (13). To date, several studies on HAI-1 and HAI-2 expression in tumor tissues have been published. The loss of

Expression of hepatocyte growth factor activator inhibitors (HAI-1 and HAI-2) in ovarian cancer

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HAI-1 expression has been implicated in the progression of breast, prostate, renal and colorectal carcinoma in humans (14-17). The matriptase to HAI-1 ratio could be important in the development of advance disease on ovarian cancer (18). One study concluded that HAI-2 expression was up-regulated in pancreatic cancers (9), whereas another demonstrated down-regulation in glioblastoma, hepatocellular carcinomas and renal carcinoma (10,19,20).

To our knowledge, HAI-1 and HAI-2 have not been reported in the same patient with ovarian cancer. In the current study, we examined whether HAI-1 and HAI-2 protein expression correlate with clinicopathological characteristics in patients suffering from ovarian cancer. The main aim of the study was to determine whether HAI-1 and HAI-2 protein could be used as an early invasive marker for patients and considered a therapeutic target for treatment approaches in ovarian cancer.

Materials and methods

Patients and tissues. Patients with malignant epithelial ovarian cancer (n=54) were treated at Okayama University Hospital. All specimens were obtained at the time of surgery from the primary lesion. Informed consent was obtained from each patient before sample collection. The histological malignant cell types of tumors were assigned according to the WHO classification: 32 were classified as serous cell type, 10 as endometrioid cell type, 2 as mucinous cell type and 10 as clear cell type. Surgical staging was reviewed based on the International Federation of Obstetrics and Gynecology (FIGO) staging system: 17 were stage I, 5 were stage II, 28 were stage III and 4 were stage IV. Overall survival was defined as the interval from the initial surgery to death or to March 30, 2006. The median duration of follow-up was 48 months (range 6-146). Tumor specimens were fixed in 10% neural-buffered formalin immediately after excision, dehydrated through graded series of ethanol and xylene, embedded in paraffin.

Immunohistochemistry and staining evaluation. Formalin-fixed, paraffin-embedded sections, 4 μm thick, were deparaffinized with xylene and rehydrated in ethanol. Endogenous peroxidase activity was quenched by methanol containing 0.3% hydrogen peroxidase for 15 min. Then, sections were incubated at room temperature with a HAI-1 and HAI-2 (Santa Cruz Biotechnology, Santa Cruz, CA) followed by staining using a streptavidin-biotin-peroxidase kit (Nichirei, Tokyo, Japan). The sections were counterstained with hematoxylin. The level of HAI-1 or HAI-2 staining in epithelial cells was classified into three groups by scoring the percentage of positive cells: strong (2); >50% of cells stained, moderate (1); 10-50% of cells stained, and weak (0); <10% of cells stained. Microscopic analyses were independently conducted by two examiners with no prior knowledge of the clinical data. Final decisions in controversial cases were made using a conference microscope.

Cell culture, media and generation of transfectants. OVCAR-3 (ATCC Number: HTB-161) cancer cell lines were derived from human ovarian carcinomas. OVCAR-3 ovarian cancer cells were maintained in Dulbecco’s modified Eagle’s medium (DMEM) with supplemented with 10% charcoal-stripped fetal bovine serum (FBS). Cells were maintained at 37˚C in an atmosphere with 5% CO2, in air. A cDNA encoding the whole coding region of HAI-1 or HAI-2/PB was constructed by polymerase chain reaction using full-length HAI-1 cDNA or HAI-2 cDNA as a template. Primers were used: 5'-TTGGAATTCCTAGGATGCCCCTGT CGAGGAC-3' and 5'-TTAGTCGACTCAAGGGGGCGG GTGTGGT-3' for HAI-1, and 5'-AGCTCTAGAGCCA TGCCGCACTGTGCGG-3' and 5'-TTAGTCGACTCA CAGGACCATATGTGTTCCTT-3' for HAI-2/PB. The polymerase chain reaction products were subcloned into the EcoRI/Sall site (HAI-1) or Xbal/Sall site (HAI-2/PB) of an expression plasmid pCIneo (Promega, Madison, WI). HAI-1 and HAI-2 cDNA expression vector was transient transfected into OVCAR-3 ovarian cancer cells using TransFast transfection reagent (Promega). Mock transfected cells served as a control.

Western blot analysis. Cell lysates were collected and estimated using a Protein Assay system (Bio-Rad, Hercules, CA) according to the manufacturer’s protocols. Proteins from each cell line were subjected to SDS-PAGE, and were transferred onto a nitrocellulose membrane. Polyclonal and monoclonal antibodies used for immunoblotting were as follows: HAI-1, HAI-2, Matriptase, Bcl-2, Bak (Santa Cruz Biotechnology), Hepsin (Cayman Chemical, Ann Arbor, MI), and β-actin (Sigma Chemical, St. Louis, MO). Working dilution for the primary antibodies was 1:1000. Membranes were then incubated with appropriate secondary antibodies. Expressions of antigen-antibody complexes were detected with an enhanced chemiluminesence kit (Amersham Biosciences, Piscataway, NJ).

Cell viability assay. To examine the cell viability by transient transfection with HAI-1 or HAI-2 in OVCAR-3 cells SYTO 10 green fluorescent nucleic acid stain and Dead red (ethidium homodimer-2) nucleic acid stain (Live/Dead® reduced biohazard viability/cytotoxicity kit; Invitrogen, Eugene, OR) were added to each well and incubated for 15 min. Cell fluorescence was observed by a fluorescent microscope (Olympus, Tokyo, Japan).

Matrigel invasion assay. To investigate differences in invasive ability between cells expressing HAI-1 or HAI-2, we used BD BioCoat Matrigel Invasion Chamber (BD Bioscience, Bedford, MA). Either with or without transient transfection with HAI-1 or HAI-2 in OVCAR-3 ovarian cancer cells were added in situ with 10 μg/ml of DI (Invitrogen, Carlsbad, CA) in DMEM with 10% FBS. Containing 10% FBS for 1 h. Cells (5x104) of each genotype were added to inserts, and 0.75 ml of medium was added to the bottom of each well. After 12, 24, 36 and 48 h of incubation, membranes were removed from the insert and mounted on slides, and then invading cells were counted under the microscope. Matrigel assays were performed in triplicate.

MTS assay and FITC-conjugated Annexin V. In order to evaluate the cell proliferation effects of HAI-1 and HAI-2,
MTS assay (Promega) was performed. Cells were seeded into 96-well plates to obtain a density of 5x10^4 cells/well and transfected. Cells were treated either with or without transient transfection with HAI-1 or HAI-2 treatment for 12, 24, 36 and 48 h and then MTS was added for 2 h. The absorbance was read at a wavelength of 490 nm using an ELISA plate-reader (Bio-Rad systems). Moreover, apoptosis was measured by staining with FITC-conjugated Annexin V using a MEB-Cyto Apoptosis Kit (MBL, Nagoya, Japan) according to the manufacturer’s recommendations with flow cytometric analysis.

Cell growth in monolayers and in soft agar. In monolayers of cell growth, cells were plated at a density of 3x10^4 cells/well in 6-well plates containing DMEM with 10% FBS. Cell proliferations were assessed by counting the cell numbers in triplicate after 2, 4, 6, 8 and 10 days using a hemocytometer. In soft agar, a cell suspension (1x10^4 cells/well) in 1 ml 0.2% Noble agar DMEM with 10% FBS was overlaid onto a 35 mm dish containing a 0.5%-agar base. Colonies >0.2 mm in diameter were counted on the 21st day after culture. Soft agar assays were performed in triplicate.

Statistical analysis. Statistical analyses were performed using the Mann-Whitney U test for comparison with the control and one-factor ANOVA followed by Fisher's protected least significance difference test for all pairwise comparisons. The analyses were performed with the software package StatView version 5.0 (Abacus Concepts, Berkeley, CA). Differences were considered significant at p<0.05.

Results

Expression of HAI-1 and HAI-2 were examined in human ovarian tissues by immunostaining. Expression of HAI-1 and HAI-2 were examined in human epithelial ovarian cancer tissues by immunostaining; Fig. 1 illustrates representative immunostaining patterns of HAI-1 and HAI-2. Weak epithelial staining was observed respectively in 9 cases (16.6%) and 15 cases (27.8%), moderate staining in 30 cases (55.6%) and 20 cases (37.0%), and strong staining in 15 cases (27.8%) and 19 cases (35.2%) of HAI-1 and HAI-2.

The biological parameters examined by immunostaining patterns of HAI-1 and HAI-2. Table I shows the distribution of cases scored as positive for each of the biological parameters examined, according to clinicopathological characteristics in the overall population. As expected, the expression of HAI-2 has a statistically significant association with clinicopathological parameters such as stage (p=0.031), amount of ascites (p=0.002), diameter of residual tumor size (p=0.034). The expression of HAI-1 shows a statistically significant association only with stage (p=0.040) (Mann-Whitney U test, p<0.05) (Table I).

The disease-free and overall survival analysis. HAI-1 and HAI-2 were significant in the disease-free survival (DFS) and overall survival (OS) analysis of prognostic factor using the log-rank test in ovarian cancer. Fig. 2 shows DFS and OS curves of 54 patients with ovarian cancer, according to HAI-1 and HAI-2 expression status. The DFS and OS rates...
of patients exhibiting high HAI-1 and HAI-2 expression (score 2) were significantly higher than those of patients exhibiting low HAI-1 and HAI-2 expression (score 0-1). (p=0.031 and p=0.003, p=0.048 and p=0.001, respectively) (Mann-Whitney U test, p<0.05).

The protein expression of HAI-1, HAI-2, hepsin and matriptase by transient transfection with HAI-1 or HAI-2 into OVCAR-3 ovarian cell lines. We examined the functional role of HAI-1 and HAI-2 which were generated as described in Materials and methods. As shown in Fig. 3A, HAI-1 and HAI-2 levels were almost absent in the OVCAR-3 cancer celllines. HAI-1 and HAI-2 protein levels were highly expressed in the transient transfection with HAI-1 and HAI-2. The expression of matriptase and hepsin was decreased by transient transfection with both HAI-1 and HAI-2. Furthermore, matriptase was inhibited more than hepsin by HAI-1 and HAI-2.

Cell viability by transient transfection with HAI-1 or HAI-2 into OVCAR-3 ovarian cancer cell lines. We investigated the efficacy of cell viability of OVCAR-3 ovarian cancer cells by transient transfection with HAI-1 or HAI-2. The percentage of cell viability was decreased to 35.6% and 48.8% of control at 48 h after transient transfection with HAI-1 or HAI-2 into OVCAR-3 cell lines, respectively. Therefore, the dead cells were increased by transient transfection with HAI-1 and HAI-2 into OVCAR-3 cells (Fig. 3B).

Matrigel invasion of transient transfection with HAI-1 or HAI-2 into OVCAR-3 cells. We assessed motility and invasiveness of cells overexpressing HAI-1 and HAI-2 using a BioCoat Matrigel Invasion Chamber. Cells were plated on the upper surface of the membrane, cells on the bottom of the membrane were stained and counted. The percentage of cells reaching the bottom of the filter was decreased to 90.6%, 86.6% of control at 12 h, 64.2%, 58.4% at 24 h, 37.6%, 32.2% at 36 h, and 29.8%, 25.5% at 48 h after transient transfection of HAI-1 or HAI-2 into OVCAR-3 cells, respectively (Fig. 4A).

Apoptosis of the transient transfection with HAI-1 or HAI-2 into OVCAR-3 cell lines. We used MTS and FITC-conjugated Annexin V assays that examined apoptosis expression by transient transfection of HAI-1 or HAI-2 into OVCAR-3 ovarian cell lines. The transient transfection of HAI-1 and HAI-2 caused reduced cell growth in MTS assay (Fig. 4B). Representative flow cytometric data revealed that transient

Table I. Association between HAI-1 or HAI-2 and clinicopathological factors in ovarian cancer.

<table>
<thead>
<tr>
<th>Variable</th>
<th>No. of cases (n=54)</th>
<th>HAI-1 (mean ± SE)</th>
<th>P-valuea</th>
<th>HAI-2 (mean ± SE)</th>
<th>P-valuea</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age (years)</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>≤60</td>
<td>39</td>
<td>1.05±0.69</td>
<td>0.186</td>
<td>1.08±0.81</td>
<td>0.836</td>
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<tr>
<td>&gt;60</td>
<td>15</td>
<td>1.27±0.59</td>
<td></td>
<td>1.13±0.74</td>
<td></td>
</tr>
<tr>
<td>Stage</td>
<td></td>
<td></td>
<td>0.040*</td>
<td></td>
<td>0.031*</td>
</tr>
<tr>
<td>I + II</td>
<td>21</td>
<td>1.35±0.67</td>
<td></td>
<td>1.38±0.74</td>
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</tr>
<tr>
<td>III + IV</td>
<td>33</td>
<td>0.97±0.63</td>
<td></td>
<td>0.91±0.77</td>
<td></td>
</tr>
<tr>
<td>Grade</td>
<td></td>
<td></td>
<td>0.262</td>
<td></td>
<td>0.276</td>
</tr>
<tr>
<td>1+2</td>
<td>21</td>
<td>1.24±0.54</td>
<td></td>
<td>1.24±0.83</td>
<td></td>
</tr>
<tr>
<td>3/NE</td>
<td>33</td>
<td>1.03±0.73</td>
<td></td>
<td>1.00±0.75</td>
<td></td>
</tr>
<tr>
<td>Histological type</td>
<td></td>
<td></td>
<td>0.139</td>
<td></td>
<td>0.495</td>
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<tr>
<td>Non-serous</td>
<td>22</td>
<td>1.27±0.63</td>
<td></td>
<td>1.18±0.80</td>
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</tr>
<tr>
<td>Serous</td>
<td>32</td>
<td>1.00±0.66</td>
<td></td>
<td>1.03±0.78</td>
<td></td>
</tr>
<tr>
<td>Amount of ascites (ml)</td>
<td></td>
<td></td>
<td>0.672</td>
<td></td>
<td>0.002*</td>
</tr>
<tr>
<td>&lt;1000</td>
<td>34</td>
<td>1.14±0.67</td>
<td></td>
<td>1.30±0.74</td>
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</tr>
<tr>
<td>≥1000</td>
<td>20</td>
<td>1.06±0.66</td>
<td></td>
<td>0.65±0.70</td>
<td></td>
</tr>
<tr>
<td>Diameter of residual tumor size</td>
<td></td>
<td></td>
<td>0.639</td>
<td></td>
<td>0.034*</td>
</tr>
<tr>
<td>≤2 cm</td>
<td>35</td>
<td>1.14±0.65</td>
<td></td>
<td>1.26±0.74</td>
<td></td>
</tr>
<tr>
<td>&gt;2 cm</td>
<td>19</td>
<td>1.05±0.71</td>
<td></td>
<td>0.79±0.79</td>
<td></td>
</tr>
<tr>
<td>Lymph node metastasisb</td>
<td></td>
<td></td>
<td>0.164</td>
<td></td>
<td>0.499</td>
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<tr>
<td>Negative</td>
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<td>1.21±0.70</td>
<td></td>
<td>1.15±0.80</td>
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</tr>
<tr>
<td>Positive</td>
<td>21</td>
<td>0.95±0.59</td>
<td></td>
<td>1.00±0.77</td>
<td></td>
</tr>
</tbody>
</table>

*aMann-Whitney U test. bLymph node status was assessed by CT imaging in 23 patients.
Figure 2. Kaplan-Meier plots for (A) disease-free and (B) overall survival of the 54 patients with ovarian cancer according to their epithelial HAI-1 expression status. Kaplan-Meier plots for (C) disease-free and (D) overall survival of the 54 patients with ovarian cancer according to their epithelial HAI-2 expression status. Low epithelial expression, score 0-1; high epithelial expression, score 2.

Figure 3. Evaluation of transient transfection of HAI-1 and HAI-2 into OVCAR-3 ovarian cancer cells. (A) The transient transfection of HAI-1 or HAI-2 into OVCAR-3 ovarian cancer cells were analyzed for HAI-1, HAI-2, matriptase, and hepsin expression by Western blotting. β-actin antibody was used as loading control in the same blot. (B) The cell viability was examined by transient transfection with HAI-1 or HAI-2 into OVCAR-3 ovarian cancer cell lines. Cell viability was analyzed by a fluorescent microscope.
transfection of HAI-1 and HAI-2 for 48 h showed increased FITC positive and Propidium iodide (PI) positive signal. HAI-1 and HAI-2 induced apoptosis in these cancer cells, which was assessed by Annexin V-FITC assay as shown in Fig. 4C. HAI-1 and HAI-2 respectively increased the total apoptotic and necrotic cells to 43.43% and 62.88% as compared with control of 6.92% in OVCAR-3 cells.

The protein expression of Bcl-2 and Bak by transient transfection with HAI-1 or HAI-2 into OVCAR-3 cell lines. We also investigated the effect of HAI-1 and HAI-2 on apoptotic molecules such as anti-apoptotic Bcl-2 and pro-apoptotic Bak on the protein level. As shown in Fig. 5A, the expression of Bak protein was up-regulated, while Bcl-2 was down-regulated in the transient transfection with HAI-1 and HAI-2 in OVCAR-3 cell lines.

Inhibition of cell growth in a monolayer and in soft agar by transient transfection of HAI-1 or HAI-2 into OVCAR-3 cell lines. Effects of HAI-1 or HAI-2 expression on cell proliferation were analyzed using transient transfection of HAI-1 or HAI-2 into OVCAR-3 cell lines. We found significant inhibitory effect of HAI-1 or HAI-2 on cell growth in transient transfection of HAI-1 or HAI-2 into OVCAR-3 cell lines (p<0.05) as compared to the control (Fig. 5B). Furthermore, the transient transfected with HAI-1 or HAI-2 into OVCAR-3 cell lines resulted in a significantly reduced number of colonies in the colony formation assay when compared to the control (p<0.01) (Fig. 5C). Furthermore, we found a significant inhibitory effect of HAI-2 compared with HAI-1 on ovarian cancer.

Discussion

HAI-1 and HAI-2 are a Kunitz-type serine protease inhibitor that has a broad inhibitory spectrum against serine proteases, such as plasmin, trypsin, tissue, and plasma kallikreins and factor XIa. cDNA cloning of HAI-2 revealed that this protein was identical to placental bikunin (PB) (21). To date, several studies on HAI-1 and HAI-2 expression in ovarian cancer tissues have been published. The matriptase to HAI-1 ratio could be important in the development of advance stage in ovarian cancer (18). Tanaka and colleagues have reported that lowest HAI-2/PB expression may be associated with
advanced stages, positive nodal status, and positive peritoneal status, and low HAI-2/PB expression correlates with poor prognosis in ovarian cancer (22). In the current study, we examined whether HAI-1 and HAI-2 protein expression correlated with clinicopathological characteristics in patients suffering from ovarian cancer by immunohistochemistry. Immunohistochemistry on ovarian cancer tissues showed that HAI-1 expression associated with stage (p=0.040). There was a significant correlation between HAI-2 expression and stage (p=0.031), amount of ascites (p=0.002) and diameter of residual tumor size (p=0.034) (Table I). Moreover, the low HAI-1 and HAI-2 expression was a significant predictor for poor prognosis compared with high HAI-1 and HAI-2 expression (DFS and OS rates: p=0.031, p=0.003 and p=0.048, p=0.001) (Fig. 2).

HAI-1 and HAI-2 are endogenous inhibitors of matriptase and hepsin. Both matriptase and hepsin are type II transmembrane proteins with an extracellular serine proteinase domain, and show enhanced expression in a variety of tumor tissues (23). Matriptase has been proposed to initiate signaling and proteolytic cascades through its ability to activate plasminogen activators such as uPA and plasminogen activators in vitro, and some in vivo evidence exists to support the physiologic role of matriptase in activating these proteins (24,25), and hepsin is thought to be involved in the activation of the coagulation cascade such as blood clotting factors VII, XII, and IX, and activate pro-uPA, and pro-HGF (26). Generation of plasmin from plasminogen by uPA can induce extracellular matrix degradation and promote tumor cell migration and metastasis. In this study, we investigated the mechanism that correlated between HAI-1, HAI-2, matriptase and hepsin in ovarian cancer cell lines. Both matriptase and hepsin were significantly inhibited by HAI-1 and HAI-2 (Fig. 3A). However, the biological functions of HAI-1 and HAI-2 remain unknown. Previous reports also showed that HAI-2/PB acts during apoptosis (27,28). In this study, we used MTS and FITC-conjugated Annexin V assays that examined apoptosis expression by transient transfection of
HAI-1 or HAI-2 into OVCAR-3 ovarian cancer cells. HAI-1 and HAI-2 induced apoptosis on OVCAR-3 ovarian cancer cells, as shown in Fig. 4B and C. To trace the steps in the apoptotic cascade, we evaluated the activation of the pro-apoptotic and anti-apoptotic proteins (29,30). Our results indicated that HAI-1 and HAI-2 up-regulated the Bak proteins and down-regulated the Bcl-2 thereby inducing apoptosis (Fig. 5A). Overall, HAI-1 and HAI-2 show potential inhibitory effects mediated by reduction of matrintsase and hepsin expression which leads to apoptosis through increasing the level of Bak and reducing the level of Bcl-2 in ovarian cancer. Therefore, we found a significant inhibitory effect of HAI-2 compared with HAI-1 on ovarian cancer in vitro and in vivo.

In summary, in vitro and in vivo studies revealed a critical role for HAI-1 and HAI-2 in disruption of the basement membrane of ovarian cancer. These findings identify HAI-1 and HAI-2 as a favorable prognostic marker, and could be considered a therapeutic target for treatment approaches in ovarian cancer.

Acknowledgments

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References


