Galectin-7 promotes the invasiveness of human oral squamous cell carcinoma cells via activation of ERK and JNK signaling

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Abstract. Galectin-7 is a member of the β-galactoside-binding protein family, and is highly expressed in oral squamous cell carcinoma (OSCC). The aim of the present study was to investigate the effects of manipulating galectin-7 expression on the biological phenotype of human OSCC cells and the associated molecular mechanisms. Knockdown of endogenous galectin-7 via small interfering RNA (siRNA) was performed and cell proliferation, apoptosis, migration, and invasion were subsequently assessed. The data indicated that galectin-7 silencing had no impact on the proliferation or apoptosis of OSCC cells. However, compared with non-transfected cells, percentage wound closure was significantly lower in galectin-7-silenced cells following 24 h incubation, indicating decreased cell migration. Furthermore, Matrigel invasion assays demonstrated that galectin-7 knockdown significantly reduced the number of invaded cells, compared with the number in non-transfected cells. Western blot analysis indicated that galectin-7 overexpression resulted in a significant increase in the expression of the proteins matrix metalloproteinase (MMP)-2 and MMP-9. The invasive abilities of cells over-expressing galectin-7 significantly decreased following co-transfection with MMP-2- or MMP-9-specific siRNA. Increasing galectin-7 expression significantly enhanced the phosphorylation of extracellular signal-related kinase (ERK) 1/2 and c-Jun N-terminal kinase (JNK) 1/2. Pharmacological inhibition of ERK or JNK activity significantly suppressed the invasiveness of galectin-7-overexpressing cells and abrogated the upregulation of MMP-2 and MMP-9. Taken together, the results of the current study provide novel evidence for the pro-invasive activity of galectin-7 in OSCC cells, which is associated with activation of ERK and JNK signaling and the induction of MMP-2 and MMP-9.

Introduction

Oral squamous cell carcinoma (OSCC) is one of the most common malignancies in the world, and has a high mortality rate (1). Invasive growth via the lymphatic route is a typical feature of OSCC (2) and lymph node involvement status has been identified as a reliable prognostic indicator in OSCC patients (3,4). Therefore, a number of studies are underway in order to investigate the molecular mechanisms involved in regulating OSCC invasiveness (5).

Galectin-7 is a member of the β-galactoside-binding protein family. It is predominantly expressed in epithelial cells within healthy tissue and plays an important role in epithelial development and homeostasis (6,7). Galectin-7 expression may be altered in epithelial cancer, therefore it may serve an important role in cancer progression (8). The exact role of galectin-7 may vary in different types of cancers; it may play distinct and even opposing roles in tumor development. For example, in human gastric cancer specimens, galectin-7 is underexpressed due to epigenetic modifications and this suppresses the proliferation and invasion of gastric cancer cells (9). By contrast, in high-grade breast cancer galectin-7 is overexpressed, facilitating the spontaneous metastasis of breast cancer cells in preclinical mouse models (10). The tumor-promoting role of galectin-7 has also been noted in ovarian cancer cells (11) and cervical cancer cells (12).

Previous studies have demonstrated that galectin-7 increases the expression of matrix metalloproteinases (MMPs), especially MMP-9, thus modulating the invasiveness of cancer cells (11,12). Additionally, OSCC tissues exhibit increased MMP-2 and MMP-9 activity compared with adjacent healthy tissues (13). It has previously been demonstrated that MMPs serve a critical role in the invasion and metastasis of oral cancer (14). Alves et al (15) reported that galectin-7 is highly expressed in OSCC and its expression is significantly correlated with the histological grade of disease. These findings suggest that galectin-7 may contribute to OSCC invasiveness by modulating the expression of MMP-2 and MMP-9. The present study investigated the effects of manipulating galectin-7 expression on the biological phenotypes of human...
OSCC cells and evaluated the involvement of MMP-2 and MMP-9 on the action of galectin-7.

Materials and methods

Cell culture and treatment. The human OSCC cell lines SCC-4 and SCC-9 were purchased from the American Type Culture Collection (ATCC; Manassas, VA, USA). All cells were maintained at 37°C in 5% CO₂ in Dulbecco’s Modified Eagle’s Medium supplemented with 10% fetal bovine serum (FBS), 1 mmol/L L-glutamine, and 100 U/ml penicillin, 100 µg/ml streptomycin (all from Invitrogen; Thermo Fisher Scientific, Inc., Waltham, MA, USA). For inhibitor experiments, cells were pretreated with the c-Jun N-terminal kinase (JNK) inhibitor SP600125 (10 µM; Calbiochem; EMD Millipore, Billerica, MA, USA), extracellular signal-related kinase (ERK) inhibitor PD98059 (10 µM; Calbiochem; EMD Millipore), or 0.1% dimethyl sulfoxide (DMSO) used as vehicle control 1 h before transfection of galectin-7-expressing plasmid.

Plasmids, small interfering RNA (siRNA), and transfection. A galectin-7-expressing plasmid (pCEP4-GAL7) was purchased from Addgene (Cambridge, MA, USA) and an empty vector (pCEP4) was also purchased (Invitrogen; Thermo Fisher Scientific, Inc.) Galectin-7 siRNA, MMP-2 siRNA, MMP-9 siRNA, and negative control siRNA were obtained from Santa Cruz Biotechnology (Dallas, TX, USA). For overexpression or knockdown of galectin-7, cells were seeded onto 6-well plates (4x10⁴ cells/well) and transfected with 1 µg pCEP4-GAL7, 1 µg pCEP4 and 50 nM galectin-7 siRNA, or 50 nM control siRNA using Lipofectamine® 2000 (Invitrogen; Thermo Fisher Scientific, Inc.), according to the manufacturer’s instructions. Cells were incubated for 24 h, and subsequently collected for further experiments. To validate the involvement of MMP-2 and MMP-9, cells were co-transfected with 1 µg pCEP4-GAL7 and 50 nM MMP-2 siRNA, MMP-9 siRNA, or control siRNA, and tested for invasive ability following incubation for 24 h.

Cell proliferation assay. Cell proliferation was measured using the MTT assay. Transfected cells were detached and re-seeded onto 96-well plates (2x10³ cells/well). Following incubation for 1, 3, and 5 days, 0.5 mg/ml MTT (Sigma-Aldrich; Merck Millipore, Darmstadt, Germany) was added to the culture and incubated for additional 4 h at 37°C. Formazan crystals were dissolved in DMSO. Absorbance was measured at 570 nm using a multi-plate reader.

Apoptosis detection assay. Apoptosis analysis was performed using the Annexin V-FITC Apoptosis Detection kit (Nanjing KeyGen Biotech Co., Nanjing, China), according to the manufacturer’s instructions. In brief, cells were incubated with a staining solution containing fluorescein isothiocyanate (FITC)-conjugated annexin-V and propidium iodide (PI) for 10 min at 4°C in the dark. The percentage of apoptotic cells was determined using a FACScan flow cytometer with the CellQuest software (BD Biosciences, San Jose, CA, USA).

Wound-healing assay. Cells were seeded onto 6-well plates and allowed to grow to ~95% confluence. A wound was made in the monolayer using a 100-µl pipette tip. The culture was washed to remove cellular debris and incubated for 24 h at 37°C. Cells were imaged using a phase contrast microscope at different time points. The extent of wound closure was quantified by measuring its area before migration, and 24 h after migration. Results were expressed as percentage of wound closure.

Transwell invasion assay. Invasion assays were performed using Transwell chambers, which were coated with Matrigel (BD Biosciences) 24 h prior to use. The cells were subsequently harvested and resuspended in serum-free medium containing 1% bovine serum albumin (Sigma-Aldrich; Merck Millipore). The cell suspension was added to the upper chamber and the lower chamber was filled with culture medium containing 10% FBS. After incubation for 24 h at 37°C, cells on the upper surface of the chamber were removed using a cotton swab. Invaded cells on the lower surface were fixed in 4% formaldehyde, stained with 0.5% crystal violet, and counted under a microscope.

Reverse transcription-quantitative polymerase chain reaction (RT-qPCR) analysis. Total RNA was extracted from cells using the TRIzol reagent following the manufacturer's instructions (Invitrogen; Thermo Fisher Scientific, Inc.). Reverse transcription was performed using the PrimeScript First Strand cDNA Synthesis kit (Takara Biotechnology Co., Dalian, China). RT-qPCR was performed on an ABI 7500 Fast Real-Time PCR System (Applied Biosystems; Thermo Fisher Scientific, Inc.) with SYBR-Green detection mix (Takara Biotechnology Co.). The following primers were used in the current study: Galectin-7 forward 5'-TTGCTCTCCTTGCTGTGAAGACCCAC-3', and reverse 5'-AGGTTCCATGTAACCTGCTTGCA-3'; and reverse 5'-AGGTTCCATGTAACCTGCTTGCA-3'; and reverse 5'-AGGTTCCATGTAACCTGCTTGCA-3'; and reverse 5'-AGGTTCCATGTAACCTGCTTGCA-3'; and reverse 5'-AGGTTCCATGTAACCTGCTTGCA-3'; and reverse 5'-AGGTTCCATGTAACCTGCTTGCA-3'; and reverse 5'-AGGTTCCATGTAACCTGCTTGCA-3'; and reverse 5'-AGGTTCCATGTAACCTGCTTGCA-3'. PCR conditions were as follows: 95°C for 5 min, followed by 40 cycles of 95°C for 10 sec, 64°C for 30 sec, and 72°C for 30 sec. The relative galectin-7 mRNA level was calculated using the 2⁻ΔΔCq method (17) following normalization against the level of GAPDH.

Western blot analysis. Cells were lysed in radioimmunoprecipitation assay buffer (phosphate buffer solution, 1% NP40, 0.5% sodium deoxycholate, 0.1% sodium dodecyl sulphate) containing a protease inhibitor cocktail (Cell Signaling Technology, Inc., Danvers, MA, USA) on ice for 30 min. After centrifugation at 15,000 x g for 20 min, the supernatant was collected and protein concentrations were measured using a protein assay kit (Bio-Rad Laboratories, Inc., Hercules, CA, USA). Cell lysates were separated by sodium dodecyl sulfate-polyacrylamide gel electrophoresis and transferred onto nitrocellulose membranes. Membranes were probed with the following antibodies at 1:300 dilution: Rabbit anti-galectin-7 monoclonal antibody (cat. no. ab108623), rabbit anti-MMP-2 polyclonal antibody (cat. no. ab97779), mouse anti-MMP-9 monoclonal antibody (cat. no. ab19906), rabbit anti-GAPDH monoclonal antibody (cat. no. ab9701, rabbit anti-ERK1/2 polyclonal antibody (cat. no. 9101), rabbit anti-phospho-JNK monoclonal
antibody (cat. no. 4668), rabbit anti-JNK polyclonal antibody (cat. no. 9252), rabbit anti-phospho-p38 monoclonal antibody (cat. no. 4511) and rabbit anti-p38 monoclonal antibody (cat. no. 8690; all from Cell Signaling Technology, Inc.). Horse-radish peroxidase-conjugated secondary antibodies (cat. nos. sc-2004 and sc-2005; Santa Cruz Biotechnology, Inc.) were diluted at 1:2,000 prior to use. Proteins were visualized using an enhanced chemiluminescence kit (Pierce Biotechnology, Inc., Rockford, IL, USA). The blots were quantified by densitometry with the Quantity One software (Bio-Rad Laboratories).

Statistical analysis. Data are expressed as mean ± standard deviation. Statistical differences were examined using one-way analysis of variance (ANOVA) followed by Tukey’s post hoc test. P<0.05 was considered to indicate a statistically significant difference.

Results

Galectin-7 silencing has no impact on cell proliferation or apoptosis in OSCC cells. To analyze the role of galectin-7 in the proliferation of OSCC cells, galectin-7-specific siRNA was transiently transfected into SCC-4 and SCC-9 cell lines. The delivery of galectin-7 siRNA significantly decreased mRNA and protein levels of endogenous galectin-7 in both SCC-4 and SCC-9 cells (Fig. 1A and B; P<0.05). The results of the MTT assay demonstrated that this downregulation of galectin-7 did not significantly affect SCC-4 and SCC-9 cell proliferation compared with non-transfected cells over a 5-day period (Fig. 1C). Annexin-V/PI staining analysis identified comparable percentages of apoptotic cells in non-transfected and galectin-7 siRNA-transfected cells (Fig. 1D).

Galectin-7 knockdown attenuates the migration and invasion of OSCC cells. The effect of galectin-7 downregulation on the invasive properties of OSCC cells was then analyzed. Galectin-7 silencing caused a significant decline in cell motility during in vitro wound-healing assays. Compared to non-transfected SCC-4 cells, the percentage wound closure was significantly lower in galectin-7-silenced SCC-4 cells 24 h following incubation (18.5±3.2% vs. 54.4±6.4%, P<0.05; Fig. 2A). Similarly, galectin-7 siRNA transfection resulted in a significant reduction in the motility of SCC-9 cells (P<0.05). Matrigel invasion assays demonstrated that galectin-7 knockdown significantly reduced the numbers of invaded cells by >60%, compared with non-transfected cells (P<0.05; Fig. 2B).

Overexpression of galectin-7 accelerates the migration and invasion of OSCC cells. Further tests confirmed the effect of increased galectin-7 on the migration and invasion of OSCC cells. Transfection of the plasmid pCEP4-GAL7 into SCC-4 and SCC-9 cells led to a significant increase in galectin-7 expression compared with non-transfected cells (Fig. 3A). This increase in galectin-7 expression in turn significantly increased OSCC cell migration and invasion (P<0.05; Figs. 3B and C).

Upregulation of MMP-2 and MMP-9 mediates the pro-invasive activity of galectin-7. A possible association between galectin-7-mediated invasiveness and MMP-2 and MMP-9 upregulation was investigated. Western blot analysis demonstrated that galectin-7 overexpression resulted in a 3.5-fold increase in MMP-2 protein and 2.3-fold increase in MMP-9 protein expression in both SCC-4 and SCC-9 cells transfected with control or galectin-7 siRNA. Bar graphs represent mean ± SD of three independent experiments. *P<0.05 vs. non-transfected cells. (C) The viability and (D) apoptosis of SCC-4 and SCC-9 cells transfected with control or galectin-7 siRNA were determined by MTT assay and annexin-V/PI staining analysis, respectively. GAL7, galectin-7; OSCC, oral squamous cell carcinoma; RT-qPCR, reverse transcription-quantitative polymerase chain reaction; PI, propidium iodide; siRNA, small interfering RNA.

Galectin-7 promotes OSCC cell invasion via activation of ERK and JNK signaling. Finally, the signaling pathways involved in the action of galectin-7 were investigated. As shown in Fig. 5A, increasing galectin-7 expression markedly enhanced the phosphorylation of ERK1/2 and JNK1/2
in SCC-4 and SCC-9 cells, without altering total levels of ERK1/2 and JNK1/2. No change in p38 phosphorylation levels was detected. Notably, the pharmacological inhibition of ERK or JNK activity significantly suppressed the invasiveness of galectin-7-overexpressing SCC-4 and SCC-9 cells (P<0.05; Fig. 5B) and abrogated the upregulation of MMP-2 and MMP-9 (P<0.05; Fig. 5C).

Discussion

Matsukawa et al (18) reported previously that adenoviral delivery of the galectin-7 gene may induce modest apoptosis and reduce the viability of human OSCC HSC3 cells. However, knockdown of galectin-7 using antisense galectin-7 oligonucleotides demonstrated no significant effects on cell viability. In the current study, the biological roles of galectin-7 in two other OSCC cell lines were explored, and targeted reduction of galectin-7 via siRNA technology did not alter viability and spontaneous apoptosis in SCC-4 and SCC-9 cells. These results suggest that galectin-7 is not required for the maintenance of OSCC cell viability. The anti-viability effect elicited by overexpression of galectin-7 may only reflect a non-specific cytotoxicity, as the potential cytotoxic activity of galectin-7 overexpression on healthy human cells was not tested in the current study.

The ability of galectin-7 to modulate cell behavior seems to be cell-dependent. Previous studies have demonstrated that overexpressing galectin-7 inhibits the proliferation of several specific cancer cells such as gastric cancer cells (9) and colon carcinoma cells (19). However, in other cancer cells including epithelial ovarian cancer (20), galectin-7 was involved in cell proliferation, as its downregulation inhibited the proliferation of A2780-PAR ovarian cancer cell.

Metastasis is the main cause of cancer-associated mortality. Galectin-7 exhibits the ability to modulate the metastatic phenotype of several types of cancer cells (10,12,21). Demers et al (21) demonstrated that ectopic expression of
galactin-7 increases the invasiveness of lymphoma, accelerates the development of thymic lymphoma, and previously identified that overexpressing galactin-7 enhances the metastatic growth of breast cancer cells in the lungs and bones in two different mouse models (10). Enforced expression of galactin-7 also promotes the invasiveness of human HeLa cervical epithelial adenocarcinoma cells (12). The results of the present study are consistent with results from previous studies, as they demonstrated that galactin-7 has the ability to modulate the invasive properties of OSCC cells. Knockdown of galactin-7 suppressed the migration and invasion of SCC-4 and SCC-9 cells, whereas overexpressing galactin-7 increased them. Taken together, these findings indicate that galactin-7 is a potential target for the treatment of tumor dissemination in OSCC.

Compelling evidence suggests that the induction of MMPs plays a pivotal role in the OSCC invasiveness. For instance, Bedal et al (22) have previously reported that collagen XVI facilitates the invasion of OSCC cells by inducing MMP-9 expression. It has previously been suggested that the downregulation of MMP-2 and MMP-9 may account for the decreased invasiveness of OSCC cells due to the knockdown of BubR1, a critical component of spindle assembly checkpoint (23). Inhibiting MMP-2 and MMP-9 expression has also been demonstrated to mediate the anti-invasive effects of curcumin (a natural polyphenolic compound) in OSCC cells (24). In line with its pro-invasive activity, galactin-7 expression increases the expression of MMP-9 in several cancer cells (12,16,21). The current study investigated the effects of MMP-2 and MMP-9 on galactin-7 action in OSCC cells. Galactin-7 overexpression resulted in the significant upregulation of MMP-2 and MMP-9.

Most importantly, silencing MMP-2 or MMP-9 significantly impaired the invasiveness of OSCC cells that overexpressed galactin-7. Thus MMP-2 and MMP-9 may be required for the galactin-7-mediated invasiveness of OSCC cells.

To gain a better insight into the function of galactin-7 in OSCC cell invasiveness, the signaling pathways involved were analyzed. Since mitogen-activated protein kinase (MAPK) pathways are implicated in the invasion of oral cancer cells (25,26) and galactin-7 may activate p38 MAPK signaling in cervical cancer cells (12), the current study investigated the importance of MAPK signaling in mediating galactin-7 action. The results of the present study demonstrated that galactin-7 overexpression leads to the phosphorylation and activation of ERKs and JNKs, but not p38 MAPK, in SCC-4

Figure 4. Upregulation of MMP-2 and MMP-9 mediates the pro-invasive activity of GAL7. (A) Western blot analysis of MMP-2 and MMP-9 protein expression in cells transfected with vector or GAL7-expressing plasmid. Bar graphs represent quantitative data from three independent experiments. *p<0.05 vs. vector-transfected cells. (B) The invasiveness of cells transfected with indicated constructs was determined by Transwell invasion assay. *p<0.05 vs. vector-transfected cells; #p<0.05 vs. cells transfected with GAL7-expressing plasmid alone. GAL7, galectin-7; MMP, matrix metallopeptase.

Figure 5. GAL7 promotes OSCC cell invasion via activation of ERK and JNK signaling. (A) Western blot analysis of signaling proteins in cells transfected with vector or GAL7-expressing plasmid. Representative blots from three independent experiments are shown. (B and C) Cells were transfected with vector or GAL7-expressing plasmid with or without prior treatment with an ERK inhibitor (Ei) or JNK inhibitor (Ji). (B) Cell invasiveness and (C) MMP-2 and MMP-9 expression were determined by Transwell invasion assay and Western blot analysis, respectively. Bar graphs represent quantitative data from three independent experiments. *p<0.05 vs. vector-transfected cells; #p<0.05 vs. cells transfected with GAL7-expressing plasmid alone. GAL7, galectin-7; OSCC, oral squamous cell carcinoma; MMP, matrix metallopeptase; ERK, extracellular signal-related kinase; JNK, c-jun N-terminal kinase.
and SCC-9 cells. Interestingly, the pharmacological inhibition of ERK or JNK activity significantly attenuated OSCC cell invasiveness induced by galectin-7 overexpression. Moreover, galectin-7-mediated upregulation of MMP-2 and MMP-9 was compromised by pretreatment with the ERK or JNK inhibitors. Taken together, these results suggest that galectin-7 promotes the invasiveness of OSCC cells largely by inducing the expression of MMP-2 and MMP-9 via activation of ERK and JNK signaling.

In conclusion, the current study provides novel evidence demonstrating the pro-invasive activity of galectin-7, which is associated with increased MMP-2 and MMP-9 expression, in OSCC cells. Further studies are required to investigate the utility of galectin-7 as a target for the treatment of metastatic OSCC.

References