Abstract. Metastasis is the primary cause of cancer-associated mortality. The ternary IPP complex of integrin-linked kinase, PINCH and parvin functions as a signaling platform for integrins, which modulate numerous cellular processes including cell migration and invasion. Chelidonine, isolated from Chelidonium majus, is a benzophenanthridine alkaloid that exhibits anticancer properties; however, the anti-migratory and anti-invasive effects of chelidonine remain unknown. The aim of the present study was to investigate the inhibitory effects of chelidonine on migration and invasion of MDA-MB-231 human breast cancer cells, and to determine the underlying mechanisms. Chelidonine was shown to inhibit the migration and invasion of MDA-MB-231 cells in a concentration-dependent manner, without affecting the cell viability. Chelidonine did not significantly inhibit the adhesion of the cells to type I collagen (COL-I), however it did affect cell spreading and reorganization of the actin cytoskeleton. Chelidonine also inhibited COL-I-induced protein kinase B (Akt) activation and translocation to the plasma membrane, however, it did not significantly inhibit the activation of focal adhesion kinase. Notably, chelidonine treatment significantly inhibited COL-I-induced formation of the IPP complex and activation of IPP downstream signaling molecules, such as extracellular signal-regulated kinase (ERK)1/2. These results suggest that chelidonine may be a potential therapeutic agent against metastasis of invasive human cancer cells.

Introduction

Metastasis is the leading cause of morbidity and mortality associated with cancer. The development of metastasis consists of numerous processes, in which cancer cells initially detach from the primary tumor, invade the surrounding tissues and intravasate into blood and/or lymphatic systems. This is followed by extravasation from the vasculature and subsequent colonization of target organs (1). The extracellular matrix (ECM) provides both structural support and extracellular cues that regulate invasive tumor growth. Tumor-associated changes in the ECM contribute to cancer progression (2). Cell migration involves the assembly and disassembly of focal adhesions. The migration of cells is stimulated extracellularly and is initiated by integrins and intracellular signaling proteins, which are located within focal adhesions (2,3). Integrins are α and β heterodimeric cell-surface receptors that mediate cell-ECM interactions, and have important roles in the regulation of normal and tumor cell migration and survival (2,3). The binding of a ligand to the extracellular integrin domain induces conformational changes and integrin clustering, that results in the activation of signaling cascades and recruitment of multi-protein complexes to focal adhesions (4). Subsequently, integrins lacking kinase activity transmit signals through a variety of intracellular protein kinases and adaptor molecules, including focal adhesion kinase (FAK), integrin-linked kinase (ILK) and paxillin (5-7).

ILK has a central role in mediating signal transduction, initiated by cell-ECM interactions. This leads to the regulation of numerous biochemical processes, including proliferation, survival, differentiation, migration, invasion and angiogenesis (8). ILK was initially described as an integrin β1 subunit-binding protein, with involvement in kinase signaling through inhibiting the formation of the IPP complex and subsequent down-regulation of IPP downstream signaling molecules, such as Akt and ERK1/2. These results suggest that chelidonine may be a potential therapeutic agent against metastasis of invasive human cancer cells.
pathways (5). However, previous studies have demonstrated that ILK is a pseudokinase that acts as a distinct adaptor protein linking integrin and α-parvin (9,10). Structurally, ILK is comprised of an N-terminal ankyrin repeat domain and a C-terminal kinase-like domain (10,11). Through the ankyrin repeat domain, ILK binds to PINCH1 (12). The kinase-like domain of ILK interacts directly with other components of integrin-based adhesion plaques, including α-parvin (9,10,13). ILK, PINCH1, and α-parvin form the ternary complex IPP, which has emerged as an essential constituent of integrin-containing adhesion sites. IPP functions both as a structural complex, that connects integrins to the actin cytoskeleton, and as a signaling platform, that modulates numerous cellular processes (6,8). The IPP complex is a central constituent of β1 and β3 integrin-containing adhesion sites, where it regulates numerous signaling pathways, including Akt, extracellular signal-regulated kinase (ERK)1/2, and Rac1 (6,8,13).

Chelidonine is a major benzophenanthridine alkaloid derived from the plant extract of *Chelidonium majus*, which is also known as the greater celandine (family Papaveraceae), and is widely distributed in Europe and Asia. The plant extract exhibits notable antitumor and antiviral activities (14). Chelidonine is the major component of the anticancer drug Ukrain, which is a semisynthetic derivative of *C. majus* alkaloids and has antitumor activity (15). This alkaloid has been shown to induce apoptosis of primary human uveal melanoma cells (16); arrest mitosis through the inhibition of tubulin polymerization and activation of the stress-activated protein kinase/c-Jun N-terminal protein kinase (17); and down-regulate the expression of telomerase in HepG2 cells (18). The effects of chelidonine on cell migration and invasion, however, have not yet been determined. The aim of the present study was to explore whether chelidonine inhibited the migration and invasion of MDA-MB-231 human breast cancer cells, and to further elucidate the underlying mechanisms.

**Materials and methods**

**Cell culture and reagents.** MDA-MB-231 human breast cancer cells were purchased from the American Type Culture Collection (Manassas, VA, USA). The cells were maintained in RPMI-1640 medium, supplemented with 100 U/ml penicillin and 100 µg/ml streptomycin (Invitrogen Life Technologies, Carsbad, CA, USA) and 10% heat-inactivated fetal bovine serum (HyClone Laboratories, Inc., Logan, UT, USA), in a humidified 5% CO₂ atmosphere at 37°C. The chelidonine used for the present study was isolated from *C. majus* as described by previous methods (19). The chemical structure of chelidonine is shown in Fig. 1. The purity of the compound was determined to be >98%, by high performance liquid chromatography analysis. HPLC was performed using a Gilson UV/VIS 156 system (Gilson Inc., Middleton, WI, USA), in reverse phase mode on an Agilent Eclipse XDB-C18 (5 µm; 4.6 x 250 mm) column (Agilent Technologies, Santa Clara, CA, USA) at an elution rate of 1 ml/min. in the gradient 15-80% of acetonitrile in 10 mM 1-hexane sulphonic acid sodium. The chelidonine was solubilized in 100% dimethyl sulfoxide (DMSO) and used at a final concentration of <0.05% DMSO. Antibodies against phospho-Akt (Ser473), Akt, ERK1/2, phospho-ERK1/2 (Thr202/Tyr204), phospho-FAK (Tyr397), FAK, and ILK were obtained from Santa Cruz Biotechnology, (Santa-Cruz, CA, USA). Antibodies against α-tubulin and PINCH1 were from Sigma-Aldrich (St. Louis, MO, USA). Protein A/G agarose beads and antibodies against GAPDH, ErbB2, and α-parvin were obtained from Santa Cruz Biotechnology Inc. (Dallas, TX, USA). Fibronectin and type 1 collagen (COL-I) were purchased from BD Biosciences (Franklin Lakes, NJ, USA). Fluorescein isothiocyanate (FITC)-conjugated phalloidin was from Enzo Life Sciences (San Diego, CA, USA).

**Cell viability assay.** The cytotoxic activity of chelidonine was determined using an MTT-based colorimetric assay. Briefly, the cells (1x10⁴ cells/well) were seeded in 96-well plates and allowed to grow for 24 h. Chelidonine was added to the wells at the following concentrations: 0, 1, 3, 10 µm. Following an additional 24 h incubation, the MTT solution (5 mg/ml) was added and the cells were incubated for 4 h. The experiment was performed in triplicate and the cell viability was presented as a percentage of the control.

**Immunoprecipitation and western blotting.** Immunoprecipitation and western blotting were performed as previously described (20,21). Briefly, the cells were lysed in lysis buffer, containing 50 mM Tris-HCl (pH 7.4), 150 mM NaCl, 1 mM EDTA, 5 mM sodium orthovanadate, 1% NP40 and a protease inhibitor cocktail (BD Biosciences), and centrifuged at 22,000 g for 10 min at 4°C. A total of 25 µg protein per lane was separated by sodium dodecyl sulfate (SDS)-polyacrylamide gel electrophoresis and transferred onto a polyvinylidene difluoride membrane (Millipore, Bedford, MA, USA.). The membrane was blocked with 5% non-fat milk for 2 h and subsequently incubated with the corresponding primary antibody (1:1,000 dilution) overnight at 4°C. Following washing and binding of an appropriate secondary antibody (1:5,000 dilution) coupled to horseradish peroxidase for 2 h at room temperature, the signals were visualized by enhanced chemiluminescence according to the manufacturer's instructions (Animal Genetics Inc, Kyonggi-do, Korea). For the immunoprecipitation, equal quantities of cell lysate were incubated with the appropriate antibodies, followed by an incubation with protein A/G agarose beads. The immunoprecipitates were extensively
washed and the eluted precipitates were resolved, transferred and probed with the appropriate antibodies.

**Cell fractionation.** The cells were washed with phosphate-buffered saline (PBS), incubated for 20 min in hypotonic lysis buffer (50 mM Tris-HCl, pH 7.0, 1 mM EDTA, 0.1% β-mercaptoethanol, 5 mM sodium orthovanadate, protease inhibitors cocktail) and 1 ml of the protein extract was lysed using a pre-chilled Dounce homogenizer (ThomasScientific, Swedesboro, NJ, USA), with a tight-fitting pestle (15 strokes). Unbroken cells and nuclei were pelleted at 1,000 x g at 4°C for 10 min, the supernatants were further centrifuged at 21,000 x g at 4°C for 45 min. The pellets, containing the cellular membranes, were washed three times in hypotonic lysis buffer and resuspended in lysis buffer prior to western blot analysis.

**Cell migration and invasion assays.** The cell migration and invasion assays were performed using a modified Boyden chamber (8 mm pore size; Corning Costar, Cambridge, MA, USA) as described by previous methods (20,21). Briefly, the lower surface of the filters were coated with COL-I or fibronectin as a chemoattractant. The upper surface of the filter was coated with Matrigel™ (BD Biosciences) for the invasion assay, or left uncoated for the migration assay. MDA-MB-231 cells were seeded at a density of 5x10⁴ cells in 100 µl RPMI, containing 0.5% bovine serum albumin (migration) or RPMI, containing 10% FBS (invasion) in the upper compartment of transwell. The lower compartment contained 800 µl RPMI, containing 10% FBS. Following incubation for 8 h (migration) or 24 h (invasion) at 37°C in 5% CO₂, the cells which had not penetrated the filter were completely wiped away using a cotton swab and the cells which had migrated to the lower surface of the filter were fixed with methanol. The cells were subsequently stained and counted in ≥5 randomly selected microscopic fields (magnification, ×100) per filter using an Olympus CKX41 inverted microscope (Olympus, Tokyo, Japan).

**Adhesion and spreading assays.** The cell adhesion assays were carried out using 96-well tissue culture plates. The plates were coated with 10 µg/ml of COL-I overnight at 4°C. Each well was rinsed with 1X PBS and blocked with PBS supplemented with 0.1% bovine serum albumin for 1 h. The cells were plated at a density of 2x10⁴ cells/well. Following a 1 h incubation, the unbound cells were removed from the wells by gentle aspiration and washed three times with PBS. The attached cells were quantified by measuring the acid phosphatase activity. Briefly, the attached cells were treated with lysis buffer (0.1 M sodium acetate, pH 5.0, 0.1% triton X-100) containing 5 mM p-nitrophenyl phosphate and incubated for 1 h at 37°C, followed by the addition of 1 M NaOH. The absorbance was measured at 405 nm using a microplate reader.

For the spreading assay, the cells were seeded into a 24-well plate coated with 10 µg/ml of COL-I (1x10⁴ cells/well) and incubated for 1 h. Following the incubation, the cells were fixed in 3.7% formaldehyde. The proportion of spread cells was determined using a light microscope. Non-spread cells were observed as small, round cells with little or no membrane protrusions, whereas spread cells were observed as large cells with extensive visible lamellipodia. The results represent the percentage of spread cells in five randomly selected microscopic fields (magnification, ×200).

**F-actin staining and confocal microscopy.** The cells were grown on glass coverslips coated with COL-I, fixed with 4% paraformaldehyde and permeabilized in 0.2% Triton X-100. The F-actin staining was performed using fluorescein isothiocyanate (FITC)-conjugated phalloidin. Confocal images were acquired using a Zeiss LSM510 META NLO inverted laser scanning confocal microscope (Carl Zeiss AG; Oberchocken, Germany; Korea Basic Science Institute Chunchen Center) equipped with an external argon HeNe laser and HeNe laser II. The images were captured at the colony midsection using a C-Apochromat 63X NA1.2 water immersion objective (Carl Zeiss AG).

**Statistical analyses.** The data represent the means ± standard deviation of three independent experiments, each repeated in triplicate. The data were analyzed by Student’s t-test. A P<0.05 was considered to indicate a statistically significant difference, as compared with the vehicle-treated control.

**Results**

**Chelidonine inhibits COL-I-induced migration of MDA-MB-231 cells without affecting cell viability.** The Boyden chamber migration assay was performed to determine
the inhibitory effects of chelidonine on the migratory ability of MDA-MB-231 cells. The lower surface of the filters was coated with COL-I and the migrated cells were analyzed following a 12 h incubation with chelidonine. Chelidonine significantly suppressed the migration of the cells towards COL-I, in a concentration-dependent manner, with a half maximal inhibitory concentration (IC₅₀) value of 1.0±0.1 µM (Fig. 2A). The effects of chelidonine on the migration of MDA-MB-231 cells induced by fibronectin, were also investigated. Chelidonine treatment weakly inhibited the migration of the cells towards fibronectin, with an IC₅₀ value of >10 µM (Fig. 2B). These results suggest that chelidonine was more effective at suppressing COL-I-induced cell migration, as compared with fibronectin-induced cell migration. The present study also examined whether chelidonine was capable of suppressing the invasive abilities of MDA-MB-231 cells, using a Boyden chamber coated with Matrigel™. The number of cells invading the lower chamber were significantly decreased when the cells were treated with chelidonine, in a concentration-dependent manner, with an IC₅₀ value of 1.4±0.2 µM (Fig. 3A). To verify that the concentrations of chelidonine used in the experiments did not affect the cell viability, MDA-MB-231 cells were incubated with various concentrations of chelidonine for 24 h and cell viability was evaluated using an MTT assay. Chelidonine did not cause any significant toxic effects on the cells <10 µM (Fig. 3B), suggesting that chelidonine is effective in inhibiting MDA-MB-231 cell migration and invasion.

Chelidonine inhibits cell spreading and actin cytoskeleton reorganization, during adhesion to COL-I. Integrin-mediated cancer cell adhesion and spreading is an important step in cell migration and invasion (22). Therefore, the present study determined whether chelidonine prevented the adhesion of MDA-MB-231 cells to COL-I. Treatment of MDA-MB-231 cells with chelidonine slightly impaired the adhesion of cells to COL-I (Fig. 4A). The percentage of adhesion inhibition, in the cells treated with 10 µM chelidonine, was 15±2.5%, which was markedly less than that of the migratory inhibition towards COL-I. The effects of chelidonine on the spreading of MDA-MB-231 cells on COL-I were also determined. Chelidonine inhibited the spreading of cells on COL-I, in a concentration-dependent manner, with an IC₅₀ value of 2.5±0.3 µM (Fig. 4B). COL-I stimulation also induced a dramatic reorganization of the actin cytoskeleton in MDA-MB-231 cells; however, treatment of the cells with 10 µM chelidonine completely blocked the COL-I-induced reorganization of the actin cytoskeleton (Fig. 4C). These results suggest that chelidonine may inhibit migration of MDA-MB-231 cells by suppressing the reorganization of the actin cytoskeleton and cell spreading.

Chelidonine inhibits COL-I-induced activation of Akt. Integrins transmit signals through a variety of intracellular protein kinases and adaptor molecules, including FAK and ILK (9,23). Activation of FAK results in numerous cellular processes, including cell attachment, migration and invasion (24). Therefore, the effects of chelidonine on COL-I-induced FAK activation were determined. COL-I stimulation did not significantly increase the protein expression levels of phospho-FAK (Tyr397) in MDA-MB-231 cells (Fig. 5A). However, COL-I stimulation did significantly increase the protein expression levels of phospho-Akt (Ser473) within 30 min, after which it gradually returned to the basal level (Fig. 5A). These results suggest that the activation of Akt may be induced at an early stage during adhesion to COL-I, in this cancer cell line. Notably, treatment of MDA-MB-231 cells with chelidonine significantly decreased phospho-Akt (Ser473) protein expression levels (Fig. 5B), in a concentration-dependent manner (Fig. 5C). Conversely, fibronectin increased the protein expression levels of phospho-FAK (Tyr397), and chelidonine treatment did not significantly alter the fibronectin-induced phosphorylation of FAK (Fig. 5D).

Chelidonine inhibits the COL-I-induced formation of the IPP complex. The IPP complex regulates multiple integrin signaling pathways, including Akt, ERK1/2 and Rac1 small GTPase (6,23). Therefore, the effects of chelidonine on the COL-I-induced formation of the IPP complex were determined.
Figure 4. The effects of chelidonine on adhesion, spreading and actin cytoskeletal reorganization of MDA-MB-231 human breast cancer cells. (A) The cells were seeded in triplicate on type 1 collagen (COL-I)-coated 96-well plates and cultured with the indicated concentrations of chelidonine for 1 h. The adhered cells were quantified by measuring the acid phosphatase activity. The data represent the average of three independent experiments, performed in triplicate; *P<0.01, as compared with the vehicle-treated control. (B) The cells were seeded on COL-I-coated 24-well plates and cultured with the indicated concentrations of chelidonine for 1 h. The cells were fixed, visualized by phase-contrast microscopy (magnification, x200) and scored for the percentage of spread cells. The data represent the average of three independent experiments, performed in triplicate; *P<0.01, as compared with the vehicle-treated control. (C) The cells were allowed to adhere to COL-I-coated (10 µg/ml) coverslips for 1 h in the presence or absence of 10 µM chelidonine, and stained with fluorescein isothiocyanate-labeled phalloidin, to detect F-actin filaments.

Figure 5. The effects of chelidonine on the activation of Akt and focal adhesion kinase (FAK) in MDA-MB-231 human breast cancer cells. (A) The cells were serum-starved for 12 h and allowed to adhere to COL-I-coated dishes (10 µg/ml) for the indicated period of time. Whole cell lysates were blotted with the indicated antibodies. (B) The serum-starved cells were allowed to adhere to COL-I-coated dishes (10 µg/ml), in the presence or absence of chelidonine (10 µM), for the indicated period of time. Whole cell lysates were blotted with the indicated antibodies. (C) The serum-starved cells were allowed to adhere to COL-I-coated dishes (10 µg/ml), with the indicated concentrations of chelidonine for 1 h. Whole cell lysates were used to determine the protein expression levels of phospho-Akt (Ser473) (p-Akt) by western blot analysis. (D) The serum-starved cells were allowed to adhere to fibronectin-coated dishes (10 µg/ml), in the presence or absence of chelidonine (10 µM), for the indicated period of time. Whole cell lysates were used to determine the protein expression levels of phospho-FAK (Tyr992) (p-FAK) by western blot analysis. H, hours.
using a co-immunoprecipitation assay. COL-I stimulation increased ILK association with PINCH1 and α-parvin, as well as Akt; however, chelidonine suppressed this association in a concentration-dependent manner (Fig. 6A). Membrane fractionation revealed that chelidonine significantly suppressed the COL-I-induced plasma membrane translocation of Akt (Fig. 6B), which is an essential process for Akt activation (25,26). To further confirm that chelidonine inhibited the COL-I-induced formation of the IPP complex, the effects of chelidonine on IPP downstream signaling molecules, such as ERK1/2, were evaluated. COL-I significantly increased the phosphorylation of ERK1/2 in MDA-MB-231 cells, in a time-dependent manner; however, chelidonine treatment decreased the protein expression levels of COL-I-induced phospho-ERK1/2 (Fig. 6C). These results suggest that chelidonine may exert its anti-migratory effects through interfering with the formation of the IPP complex.

**Discussion**

*Chelidonium majus* has a history in phytomedicine for the treatment of numerous diseases and health disturbances. It contains isoquinoline alkaloids, particularly protoberberine and benzophenanthridine alkaloids (27). The plant extract exhibits both antitumor and antiviral activities (14), in addition to hepatoprotective and anti-genotoxic effects in mice (28). Ukrain, a semi-synthetic derivative of *C. majus* alkaloids, has been used in therapy to treat various types of solid tumor. Previous pre-clinical and clinical investigations have established Ukrain as an anticancer drug (29). Chelidonine, the major component of Ukrain, induces apoptosis (16), mitotic arrest (17) and reduction of telomerase activity (18) in cancer cells. However, the underlying mechanisms behind the anticancer effects of chelidonine remain to be elucidated.

In the present study, the anti-migratory and anti-invasive effects of chelidonine on MDA-MB-231 human breast cancer cells were investigated. The results demonstrated that chelidonine exhibited a potent anti-migratory effect on MDA-MB-231 cells induced by COL-I, without affecting cell viability. Based on these results, the mechanism of action of chelidonine, for inhibiting COL-I-induced migration, was further explored. Chelidonine was shown to be capable of suppressing COL-I-induced migration, as well as reorganization of the actin cytoskeleton and cell spreading. Notably, chelidonine treatment suppressed the formation of the IPP complex and subsequent activation of IPP downstream signaling molecules, including Akt and ERK1/2 induced by COL-I. These results suggest that the anti-migratory mechanisms of chelidonine may be associated with inhibition of integrin signaling, by suppressing the COL-I-induced formation of the IPP complex.

Integrin-mediated signaling regulates a variety of biological processes, including cell migration, survival and proliferation. The ternary IPP complex is also a central constituent of adhesion sites, where it regulates multiple signaling pathways, including Akt, ERK1/2, and Rac1 (6,8,10). The present study demonstrated that chelidonine suppressed the COL-I-induced association of ILK with PINCH1, α-parvin and Akt. Furthermore, chelidonine suppressed COL-I-induced reorganization of the actin cytoskeleton, cell spreading and migration, as well as activation of the downstream IPP signaling.
molecules, including Akt and ERK1/2. Therefore, chelidonine may inhibit cell migration by suppressing COL-I-induced integrin signaling, through inhibiting the formation of the IPP complex in MDA-MB-231 cells. However, the detailed mechanisms by which chelidonine interferes with IPP complex formation induced by COL-I, remain to be elucidated.

Chelidonine was shown to be more effective at suppressing COL-I-induced cell migration, as compared with fibronectin-mediated cell migration. The binding of integrins to their ligands induces conformational changes and integrin clustering, resulting in the activation of signaling cascades and recruitment of multi-protein complexes to focal adhesions (30). FAK is predominantly activated in focal adhesions and is important in cell-ECM interactions that affect cell migration, proliferation and survival (4,7). Autophosphorylation of FAK at Tyr397, occurs in response to numerous stimuli, including integrin engagement. FAK phosphorylation promotes the Src homology domain 2-dependent binding of the Src family tyrosine kinases and the formation of an activated FAK-Src complex (7). FAK activation at focal adhesion sites enhances cytoskeletal reorganization, cellular adhesion and cell survival (7). Previous studies have shown that stimulation of MDA-MB-231 cells with fibronectin predominantly induces the activation of FAK (20,21). Conversely, COL-I stimulation does not significantly increase FAK phosphorylation at Tyr397, but induces the activation of Akt in MDA-MB-231 cells, suggesting that the formation of the IPP complex could be induced at an early time during adhesion to COL-I, in this cancer cell line. Chelidonine did not significantly decrease the level of FAK phosphorylation induced by fibronectin, but it did effectively suppress the COL-I-induced formation of the IPP complex, and activation of IPP downstream signaling molecules, including Akt. These results indicate that chelidonine may preferentially inhibit the IPP complex, rather than suppress FAK activation. This may explain why chelidonine was more effective at suppressing COL-I-induced cell migration, as compared with fibronectin-mediated cell migration.

The anticancer effects of chelidonine have previously been reported, however this is the first report, to the best of our knowledge, demonstrating that chelidonine inhibits migration and invasion of MDA-MB-231 cells. These effects may be achieved through the inhibition of the IPP complex and subsequent activation of IPP downstream signaling molecules, such as Akt and ERK1/2. The results of the present study also support potential, additional biological activities of chelidonine, and may provide a basis for the development of more specific cancer chemotherapeutic agents derived from natural products.

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