Research Article

Potential Therapeutic Role of Hispidulin in Gastric Cancer through Induction of Apoptosis via NAG-1 Signaling

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Gastric cancer is one of the most common malignant cancers due to poor prognoses and high mortality rates worldwide. However, an effective chemotherapeutic drug without side effects remains lacking. Saussurea involucrata (SI) Kar. et Kir., also known as snow lotus, grows in mountainous rocky habitats at 2600 m elevation in the Tian Shan and A'er Tai regions of China. The ethyl acetate extract of SI had been shown to inhibit proliferation and induce apoptosis in various tumor cells. In this study, we demonstrated that Hispidulin, active ingredients in SI, inhibits the growth of AGS gastric cancer cells. After Hispidulin treatment, NAG-1 remained highly expressed, whereas COX-2 expression was downregulated. Flow cytometric analysis indicated that Hispidulin induces G1/S phase arrest and apoptosis in a time- and concentration-dependent manner. G1/S arrest correlated with upregulated p21/WAF1 and p16 and downregulated cyclin D1 and cyclin E, independent of p53 pathway. In addition, Hispidulin can elevate Egr-1 expression and ERK1/2 activity, whereas ERK1/2 inhibitor markedly attenuated NAG-1 mediated apoptosis. Taken together, Hispidulin can efficiently activate ERK1/2 signaling followed by NAG-1 constitutive expression and trigger cell cycle arrest as well as apoptosis in cancer cell. It can be a potential compound for combination therapy of gastric cancer in the future.

1. Introduction

Gastric cancer is one of the most common causes of cancer-related mortality in China and other Asian countries [1, 2]. Surgery and chemotherapy are the standard treatment modalities for gastric cancer [3–5]. The 5-year survival of gastric cancer patients is currently estimated at approximately 30%; therefore, the development of novel treatment strategies to improve patients’ prognoses is urgently required. The majority of gastric adenocarcinomas express high levels of cyclooxygenase-2 (COX-2) [6–9]. Houghton et al. reported that angiogenesis and Helicobacter pylori infection are both associated with COX-2 expression in gastric cancer patients [10]. The knockdown of COX-2 in a SGC-7901 gastric adenocarcinoma cell line by RNA interference inhibited proliferation and induced apoptosis, indicating that suppression of COX-2 might represent an effective approach for the treatment of gastric cancer. The majority of selective COX-2 inhibitors exert pronounced side effects that limit their administration. Based on this clinical phenomena, AGS gastric adenocarcinoma cell line which constitutively expressed both COX-1 and COX-2 and additional COX-2 induced by IL-1β potentially make it an excellent model for assessing gastrointestinal toxicity of COX-2 inhibitors was an ideal model to evaluate the potential compound for adjuvant therapy in gastric cancer [11]. In this study, we observed that
Hispidulin inhibited the proliferation of AGS gastric adenocarcinoma cells; therefore, we aimed to establish if the mechanism underlying the antiproliferative effects of Hispidulin on AGS cells is the downregulation of COX-2 expression.

In our previous study, isochaihulactone markedly upregulated the expression of the nonsteroidal anti-inflammatory-drug- (NSAID-) activated gene-1 (NAG-1; also known as MIC-1, GDF-15, placental TGF-β, and PLAB). NAG-1 is a transforming growth factor-β-like secreted protein, initially characterized as a p53-regulated gene [12–14]. Overexpression of NAG-1 in breast cancer cells resulted in growth arrest and apoptosis in vitro and in vivo. Previous studies observed similar results in colon cancer cells [15–17] and following the treatment of prostate cancer cells with purified NAG-1 [18]. These findings suggested that NAG-1 is associated with apoptosis and that the downregulation of NAG-1 expression might promote tumorigenesis. In our previous investigation, we identified that isochaihulactone, a novel lignan isolated from the root of Bupleurum scorzonerifolium, induced microtubule depolymerization, cell cycle arrest, and proapoptotic activity in A549 human lung cancer cells [19]. We also observed that NSAIDs upregulated the expression of several isochaihulactone-induced genes and that isochaihulactone upregulated NAG-1 protein expression in a time-dependent manner. The upregulation of NAG-1 by isochaihulactone was associated with the upregulation of EGR-1 expression; therefore, the silencing of EGR-1 expression by siRNAs could also be associated with downregulated NAG-1 RNA and protein expression. In our analyses, the MEK1/2 inhibitor PD98059 reduced the inhibitory effects of isochaihulactone significantly; however, the p38 inhibitor SB203580 and the JNK inhibitor SP600125 did not limit isochaihulactone-induced growth inhibition. The MEK1/2 inhibitor PD98059 reduced isochaihulactone-induced upregulation of EGR-1 and NAG-1 protein expression, whereas SB203580 and SP600125 had nonsignificant effects on EGR-1 and NAG-1 expression. These data supported the concept that isochaihulactone-induced ERK1/2 activity is critical for the regulation of EGR-1 and NAG-1 expression. The induction of ERK1/2 activity and subsequent induction of EGR-1 and NAG-1 contributes to the growth inhibitory and apoptosis-promoting effects of antitumor compounds in cancer cells.

Saussurea involucrata Kar. et Kir., or the snow lotus, grows in mountainous rocky habitats at 2600 m elevation or higher in the Tian Shan and A'er Tai regions of China. Because of excessive harvesting of the wild plants for use in pharmaceutical preparations and their remarkably slow growth, the wild population of S. involucrata has depleted in recent years. S. Involutrate is currently close to extinction and, therefore, listed as a second-grade national protected wild plant in China [20, 21]. According to the theories of traditional Chinese medicine, S. Involutrate has the effects of warming the kidney, activating “yang,” expelling wind, eliminating dampness, inducing menstruation, and promoting blood circulation [22]. Hispidulin (40,5,7-trihydroxy-6-methoxyflavone) is a naturally occurring flavone in S. Involutrate [20]. Several studies have shown its potent antioxidative, antifungal, anti-inflammatory, antimutagenic, and antineoplastic properties in vitro [23–25]. A recent study identified Hispidulin as a potent ligand of the human central benzodiazepine receptor in vitro [26]. Hispidulin also acts as a partial positive allosteric modulator at GABAA receptors, penetrates the blood-brain barrier, and possesses anticonvulsant activity in the central nervous system (CNS) [27]. Lin et al. further reported that in Hispidulin-treated glioblastoma (GBM) cells, the activation of AMP-activated protein kinase (AMPK) suppressed protein synthesis, lipogenesis, and cell cycle progression. Their results suggested that Hispidulin might be useful as a chemopreventive or therapeutic agent for GBM. Subsequent observations indicated that Hispidulin is a potential modulator of CNS activity, prompting our own investigation of its antineoplastic activity against GBM [28].

In this study, we identified that Hispidulin treatment markedly upregulated NAG-1 protein expression and downregulated COX-2 protein expression significantly, in AGS gastric cancer cells. After various durations of exposure of cells to Hispidulin, the expression of EGR-1 and that of NAG-1 was upregulated in a time-dependent manner. Hispidulin treatment also increased ERK1/2 activity, and an ERK1/2 inhibitor markedly downregulated the expression of NAG-1 and the growth inhibitory effects of Hispidulin in AGS cells. These results suggested that the apoptotic effects of Hispidulin in human gastric cancer cells might be directly associated with the upregulation of NAG-1 expression through ERK1/2 activation. Our findings indicate that Hispidulin exerts therapeutic effects on human gastric cancer cells through the activation of NAG-1 through the ERK1/2 signaling pathway.

2. Materials and Methods

2.1. Preparation of Fractions. The wild plant of S. involucrata used in this study was a gift from Biopure Biotechnology (Changhua, Taiwan). Twenty grams of dried and powdered aerial parts, including flower, of S. involucrata was extracted with 100 mL of methanol three times under reflux for 2 h, respectively. The methanol extracts (SI-1) were combined, and the solvent was evaporated in vacuum to give a deep brown syrup. The syrup was resuspended in water and then partitioned successively with pentane, ethyl acetate (SI-2), and n-butanol (SI-3) to leave a water layer (SI-4). The solvents were evaporated, respectively, and the residues were used throughout this study.

2.1.1. Reverse-Phase High-Performance Liquid Chromatography (HPLC) Analysis of Flavonoids in S. involucrata. The determination of flavonoids from S. involucrata was carried out by HPLC with a photodiode detector. The HPLC system consisted of a Shimadzu LC-20AT solvent delivery system, equipped with a SPD-M20A photodiode array detector, set at 270 nm. Samples were injected with Sil-20A autosample to separate the TSK-Gel ODS-100S column. The column was maintained at an ambient temperature of 25° C. The flow rate of the system was 1.0 mL/min. The mobile phase consisted of solvent A (0.3% formic acid) and solvent B (acetonitrile). The elution profile for A was 0–10 min, with a linear gradient change of 0–5%; 10–40 min, with a linear gradient change to 55%; and maintained for another 10 min with a postrun time.
to equilibrate the column and for the baseline to return to the normal and initial working conditions.

2.2. Chemicals and Reagents. Rutin was dissolved in DMSO to a concentration of 50 mM and stored in −20°C as a master stock solution. Dimethyl sulfoxide (DMSO), 3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyl tetrazolium bromide (MTT), 2′,7′-dichlorofluorescein diacetate (H$_2$DCF-DA), Hoechst 33342, thiobarbituric acid (TBA), hydrogen peroxide (H$_2$O$_2$), trichloroacetic acid, and malondialdehyde (MDA), propidium iodide (PI), and actin antibody were purchased from Sigma-Aldrich (St. Louis, MO, USA). NuPAGE Bis-Tris Electrophoresis System (precast polyacrylamide minigel) was purchased from Invitrogen (Carlsbad, CA, USA). COX-2 antibody was purchased from Thermo scientific (Waltham, MA, USA). PARP antibodies and horseradish peroxidase-conjugated anti-mouse or anti-rabbit IgG secondary antibodies were purchased from Cell signaling (MA, USA). Polyvinylidene fluoride (PVDF) membranes, BSA protein assay kit, and Western blot chemiluminescence reagent were purchased from Amersham Biosciences (Arlington Heights, IL). Superoxide dismutase activity assay kit was purchased from biovision (Mountain View, CA). Glutathione peroxidase assay kit was purchased from Cayman Chemical (MI, USA). DNA Fragmentation Assay Kit was purchased from Clontech Laboratories (Mountain View, CA). Nonradioactive Cytotoxicity Assay was purchased from promega (Madison, WI, USA).

2.3. Cell Lines and Cell Culture. AGS human gastric adenocarcinoma cell line (ATCC, CRL-1739) were obtained from American Type Culture Collection (Manassas, VA) and propagated in culture dishes at the desired densities in RPMI 1640 medium supplemented with 10% fetal bovine serum and 1% penicillin/streptomycin. The cells were incubated at 37°C in a humidified atmosphere containing 5% CO$_2$.

2.4. Growth Inhibition Assay. The viability of the cells after treatment with various chemicals was evaluated using the MTT assay preformed in triplicate. Briefly, the cancer cells ($3 \times 10^5$) were incubated in 96-well plates containing 200 μL of the culture medium. Cells were permitted to adhere for 12–18 h then washed with phosphate-buffered saline (PBS). Solutions were always prepared fresh by dissolving 0.2% DMSO or drugs in culture medium and then were added to AGS cells. For inhibitor treatment experiments, cells were treated with 50 μM Hispidulin and preincubated for 1 h with 25 and 50 μM ERK1/2 inhibitor PD98059. After 48 h of exposure, the drug-containing medium was removed, washed with PBS, and replaced by fresh medium. The cells in each well were then incubated in culture medium with 500 μg/mL MTT for 4 h. After the medium was removed, 200 μL of DMSO and 25 μL of glycine buffer (0.1 M glycine and 0.1 M NaCl, pH 10.5) were added to each well. Absorbance at 570 nm of the maximum was detected by a PowerWave X Microplate ELISA Reader (Bio-Tek Instruments, Winooski, VT). The absorbance for DMSO-treated cells was considered as 100%. The results were determined by three independent experiments.

2.5. IC$_{50}$ Determination. MTT assay was according to the paper [29]. Briefly, the pale yellow redox indicator 3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyl tetrazolium bromide (MTT) is reduced to a dark blue end product, MTT-Formazan, by the mitochondrial dehydrogenases of living cells. MTT reduction can be measured spectrophotometrically at a wavelength of 570 nm. According to the dosage dependent curve, the MTT reduction half was indicated IC$_{50}$. In our study, Hispidulin is resolved on DMSO agent with final concentration 0.2%. Therefore, we used 0.2% DMSO as a control agent. The absorbance for DMSO treated cells without adding any drugs was considered as 100%.

2.6. Cell Cycle Analysis. The cell cycle was determined by flow cytometry with DNA staining to reveal the total amount of DNA. Approximately 5 × 10$^5$ of cells were incubated in various concentrations of Hispidulin for the indicated time. Cells were harvested by treating the cells with trypsin/EDTA. The cells were collected, washed with PBS, fixed with cold 70% ethanol overnight, and then stained with a solution containing 20 μg/mL PI and 0.1% Triton X-100 for 1 h in the dark. The cells will then pass through FACSscan flow cytometer (equipped with a 488-nm argon laser) to measure the DNA content. The data was obtained and analyzed with Cell Quest 3.0.1 (Becton Dickinson, Franklin Lakes, NJ) and ModFit LT V2.0 software.

2.7. Western Blot Analysis. Approximately 5 ×10$^6$ cells were cultured in 100 mm$^2$ dishes and then incubated in various concentration of Hispidulin for 48 h. The cells were lysed on ice with 150 mL of lysis buffer (50 mmol/L Tris-HCl, pH 7.5, 0.5 mol/L NaCl, 5 mmol/L MgCl$_2$, 0.5% nonidet P-40, 1 mmol/L phenylnisulfonyl fluoridetor, 1 mg/mL pepstatin, and 50 mg/mL leupeptin) and centrifuged at 13000 × g at 4°C for 15 min. The protein concentrations in the supernatants were quantified using a BSA Protein Assay Kit. Electrophoresis was performed on a NuPAGE Bis-Tris Electrophoresis System using 50 mg of reduced protein extract per lane. Resolved proteins were then transferred to polyvinylidene fluoride (PVDF) membranes. Filters were blocked with 5% nonfat milk overnight and probed with appropriate dilution of primary antibodies for 2 h at room temperature. Membranes were washed with three times with 0.1% Tween 100 and incubated with HRP-conjugated secondary antibody for 1 h at room temperature. All proteins were detected using Western Lightning chemiluminescence reagent plus and quantified using a densitometer.

2.8. Detection of Apoptosis. The viability of the cells after treatment with various chemicals was evaluated using an MTT assay preformed in triplicate. Briefly, the cancer cells ($5 \times 10^5$) were incubated in 96-well plates containing 200 μL of serum-containing medium. Cells were permitted to adhere for 12 to 18 h and then were washed with phosphate-buffered saline (PBS). Solutions were always prepared fresh by dissolving 0.2% DMSO or drugs in culture medium and added to AGS cells. For inhibitor treatment experiments, cells were treated with 50 μM Hispidulin and preincubated for 1 h...
with ERK1/2 inhibitor PD98059. After 24 h of exposure, the drug-containing medium was removed, washed with PBS, and replaced by fresh medium. The apoptosis was analyzed according to the method described by van Engeland et al. [30] to detect the integrity of cellular membrane and the externalization of phosphatidylserine (Cytometry 1998; 31:1–9). In brief, approximately 1 × 10⁶ cells were grown in 10 mm diameter plates. The cells were incubated in various concentrations of K8 for the indicated time and then labeled with FITC Annexin V and PI prior to harvesting. After labeling, the cells were washed with binding buffer and harvested by scraping. Cells were resuspended in binding buffer at a concentration of 2 × 10⁵ cells/mL before analysis by flow cytometry (FACScan). The data was analyzed on WinMDI V2.8 software. The percentage of cells undergoing apoptosis was determined by three independent experiments.

2.9. Caspase Activity Assay. Activity of caspase-3 was detected by using a fluorometric assay kit (Promega) according to the manufacturer's protocol. In brief, 2 × 10⁵ control or treated cells were lysed in 50 μL of cold lysis buffer and incubated in ice for 10 min. Fifty microliters of cell lysates was added to 50 μL of reaction buffer and 5 μL of fluorogenic report substrates specific for caspase-3 in a 96-well microplate. After incubation at 37°C for 1 h, the fluorescence from the cleaved C-terminal side of the aspartate residue of DEVD� amino-4-trifluoromethyl coumarin was detected by a fluorescence microplate reader (Fluoroskan Ascent; Thermo Fisher Scientific, Waltham, MA), with excitation at 400 nm and emission at 505 nm.

2.10. Statistical Analysis. The data was shown as mean with standard deviation. The statistical difference was analyzed using the Student's t-test for normally distributed values and by nonparametric Mann-Whitney U test for values of nonnormal distribution. Values of P < 0.05 were considered significant.

3. Results

3.1. Hispidulin Inhibits Human Gastric Cancer Cell Growth. To test the inhibitory effect of Hispidulin (40,5,7-trihydroxy-6-methoxyflavone) on cancer cells, we treated human gastric cancer AGS cells with Hispidulin. Twenty-four, 48, and 72 hours after treatment followed by MTT cell viability assay, we observed that Hispidulin displayed significantly higher cytotoxicity to AGS cells than other drugs such as rutin and Aspirin (Figure 1). The IC50 of Hispidulin to AGS cell was 50 μM at 48 h after treatment and 20 μM at 72 h after treatment (Figure 1(a)). On the other hand, the IC50 of rutin, a main component of Sassaurea involucrata that can attenuate the senescence [21], was over 500 μM (Figure 1(b)). Since cyclooxygenase-2 (COX-2) was often overexpressed in gastric adenocarcinomas, we also treated AGS cells with COX-2 inhibitor (celecoxib or NS-398). The results indicated that the IC50 of celecoxib and NS-398 were around 40 μM and 50 μM 48 h after treatment and 30 μM and 40 μM 72 h after treatment, respectively, (Figures 1(c)-(1(d)). However, treated AGS cells with COX-1 inhibitor, Aspirin, had less inhibitory effect on cell viability (IC50 > 1 mM) (Figure 1(e)). Taken together, Hispidulin can be more effective in gastric cancer survival inhibition than well-known COX-2 inhibitors.

3.2. Hispidulin Induces AGS Cells Apoptosis and G1/S Phase Cell Cycle Arrest. We then evaluated the association between Hispidulin-induced inhibition of cell growth and induction of apoptosis. After Hispidulin treatment, the percentage of cell apoptosis was significantly elevated in dose and time-dependent manners (Figure 2(a)). In addition, activation of caspase-3 was also significantly induced with dose-dependent phenomena (Figure 2(b)). Furthermore, caspase-3 was activated significantly 24 to 48 h after Hispidulin treatment (Figure 2(c)), and the caspase-3 mediated apoptosis was attenuated by pan caspase-3 inhibitor FMK (Figures 2(b-2(c)). These results indicated that the apoptosis effect by Hispidulin on AGS cell line was a caspase-dependent manner. The western blot analysis also more supported that Hispidulin induced apoptosis included activation of caspase-9, cleavage of caspase-3, and poly(ADP-ribose) polymerase (PARP) with in time- and dose-dependent manners (Figures 2(d-2(e)).

3.3. Hispidulin Induces AGS Cell Cycle Arrest during the GI/S Phase. To elucidate the mechanisms underlying the activities of Hispidulin, we evaluated its effects on cell cycle progression. Our results from flow cytometric analysis showed that Hispidulin treatment induced the accumulation of cells in the GI/S phase in a time-dependent manner (Figure 3(a)). This result suggested that Hispidulin might induce GI/S phase arrest, similar to fluorouracil (5-FU). Hispidulin also upregulated the expression of the GI/S regulatory proteins, including p53, p16, and p21, in time- and dose-dependent manners (Figure 3(b)). Hispidulin treatment downregulated the expression of the cell cyclin-associated proteins, including cyclin D1 and cyclin E, in AGS cells (Figure 3(b)).

3.4. Involvement of NAG-1 Activation and COX-2 Inhibition in Hispidulin-Induced Inhibition of AGS Cell Growth. The IC50 for Hispidulin was 50 μM in AGS cells (Figure 1(a)); therefore, we used the same concentration in this study's experiments. To evaluate the involvement of EGR-1 in the upregulation of NAG-1 expression induced by Hispidulin treatment in AGS cells, we used western blotting to analyze EGR-1 and NAG-1 protein expression. After various durations of exposure of cells to Hispidulin, the expression of EGR-1, as well as NAG-1, was upregulated in a time-dependent manner. EGR-1 was upregulated significantly after 6 h treatment and this effect remained until 24 h treatment. NAG-1 expression peaked after 48 h Hispidulin treatment. Hispidulin treatment markedly downregulated COX-2 expression and NF-kappa B subunit p65 expression in AGS cells (Figure 4).

3.5. The Involvement of ERK1/2 Signaling in the Hispidulin-Induced Upregulation of EGR-1 and NAG-1 Expression. To investigate the possible role of ERK1/2 in the regulation of EGR-1 and NAG-1, we treated AGS cells with Hispidulin in
Figure 1: Hispidulin can efficiently inhibit gastric cancer cell growth. (a) Cell viability after Hispidulin treatment. (b) Cell viability after rutin treatment. (c) Cell viability after COX-2 inhibitor, celecoxib, treatment. (d) Cell viability after COX-2 inhibitor, NS-398, treatment. (e) Cell viability after COX-1 inhibitor, Aspirin, treatment. Each column represents the mean ± SD (*P < 0.05; **P < 0.01; ***P < 0.001).
Figure 2: Hispidulin can induce cancer apoptosis by caspase-3 activation. (a) Cell apoptosis assay after treatment by flow cytometry apoptosis analysis. (b) Caspase-3 activity assay after treatment in various dosage. (c) Caspase-3 activity assay 6, 12, 24, and 48 h after 50 μM Hispidulin treatment. (d) The Western blot analysis for apoptosis related protein analysis after Hispidulin treatment. Expression of β-actin was used as an internal control. (e) The Western blot analysis for apoptosis related protein analysis 6, 12, 24, and 48 h after treatment. Expression of β-actin was used as an internal control. Each column represents the mean ± SD (* P < 0.05; ** P < 0.01; *** P < 0.001).
the presence and absence of the MEK1/2 inhibitor PD98059 (25 μM and 50 μM), the p38 inhibitor SB203580 (10 μM and 20 μM), or the JNK1/2 inhibitor SP600125 (10 μM and 20 μM). PD98059 reduced the growth inhibitory effects of Hispidulin in a dose-dependent manner (Figure 5(a)). We then investigated the effects of Hispidulin on MAPK activation to establish which MAPK pathways are involved in its growth inhibitory effects. Following the exposure of AGS cells to Hispidulin, we observed the upregulation of phosphor-ERK protein expression (Figure 5(b)). Using western blot analysis, we then identified that the inhibition of ERK1/2 expression by PD98059 induced the downregulation of EGR-1, NAG-1, and COX-2 protein expression, and reduced growth inhibition, in a dose-dependent manner in Hispidulin-treated AGS cells (Figures 5(c) and 5(d)). In contrast, inhibition of JNK1/2 exerted minimal effects on NAG-1 expression, whereas inhibition of p38 did not affect the expression of the 2 genes. These results suggested that the activation of the ERK1/2 signaling pathway is involved in the upregulation of EGR-1 and NAG-1 expression by
Hispidulin. Taken together, Hispidulin can efficiently inhibit cancer cell survival through apoptosis induction via ERK1/2, NAG-1 mediated pathway. Thus, it can be applied to clinically combined treatment for gastric cancer elimination (Figure 6).

4. Discussion

NSAIDs are effective chemopreventive agents for various cancers via the inhibition effect on prostaglandin synthesis. Previous studies identified the chemopreventive and antitumorigenic activities of NSAIDs against colorectal and other human cancers; however, the molecular mechanisms responsible for these properties have yet to be fully elucidated [31–33]. It has been reported that NAG-1 is a target gene for NSAIDs and a unique member of the transforming growth factor superfamily. Increases in NAG-1 expression result in the induction of apoptosis in several cancer cell lines [14, 16]. Also, NAG-1 expression is induced not only by NSAIDs but also by several antitumorigenic compounds. These include dietary compounds, peroxisome proliferator-activated receptor-γ ligands, and phytochemicals [34, 35] as well as resveratrol, genistein, diallyl disulfide, 5F203, and retinoid 6-[3-(1-adamantyl)-4-hydroxyphenyl]-2-naphthalene carboxylic acid [17, 36, 37]. In our previous study, after treatment with [3-(1-adamantyl)-4-hydroxyphenyl]-2-naphthalene carboxylic acid [17, 36, 37]. In our previous study, we also found that a novel lignin, isochaihulactone, increased NAG-1 mRNA and protein expression and inhibited cell proliferation on human lung cancer A549 cells. In this study, we reported that the flavonoid Hispidulin upregulates the expression of the proapoptotic and antitumorigenic protein NAG-1. Our results indicated that NAG-1 is an important target gene for Hispidulin. Our study results could increase understanding of the mechanisms by which Hispidulin can affect tumor development. During our analyses, we observed the Hispidulin-induced upregulation of NAG-1 expression in the human AGS gastric cancer cell line had relatively reduced COX-2 activity repression. Our results demonstrated that Hispidulin-treated cells had NAG-1 elevated expression and reduced COX-2 expression. However, the regulatory relationship between NAG-1 and COX-2 still remained to be investigated although many anticancer drugs and compounds had effects on NAG-1 and COX-2 at the same treatment [20, 28]. Since COX-2 overexpression had relative malignance in clinical gastric cancer patients, this finding provide a potency that Hispidulin can be served as an adjuvant therapy.

Expression of iNOS and COX-2 is largely regulated by transcriptional activation. Among these transcription factors, NF-κB, which is a primary transcription factor and regulates various genes, is critical in the inflammation [38]. NF-κB is a redox-sensitive transcription factor that regulates a multitude of inflammatory genes, including cytokines, chemokines, adhesion molecules, and acute phase proteins. Under basal conditions, NF-κB is inactive and prevented from DNA binding and nuclear translocation by tight association in the cytoplasm with inhibitory proteins. Cell activation by a variety of extracellular signals, such as oxidative stress, induces a cascade of events that lead to activating NF-κB then translocating it to the nucleus where it binds to DNA elements in the promoters of numerous proinflammatory gene families [21, 39]. In our study, after treatment with hispidulin, the expression of nuclear factor κB (NF-κB)-p65 in cytoplasm extract fractions decreased, as compared to that of the control (Figure 4). Taken together, the results suggest that hispidulin may exert anti-inflammatory effects in vitro in AGS cells through inhibition of NF-κB signal pathway activation. Induction of EGR-1 expression by anti-tumorigenic compounds is known to involve members of the family of mitogen-activated protein kinases (MAPKs) or phosphatidylinositol-3-kinase (PI3 K) dependent pathways. For example, induction of EGR-1 expression by the peroxisome proliferator-γ activated receptor-γ (PPARγ) ligand troglitazone occurs by the ERK phosphorylation pathway rather than by the PPARγ pathway [15, 16].

To determine which MAPK family is involved in the major signaling pathway for Hispidulin-mediated NAG-1
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Figure 5: Inhibition of NAG-1 expression and growth inhibition by ERK1/2 inhibitor. (a) AGS cells were treated with 30 μM Hispidulin in the presence or absence of the mitogen-activated protein kinase 1/2 inhibitor PD98059, p38 inhibitor SB203580, and JNK1/2 inhibitor SP600125. For 48 h incubation, growth inhibition effect was determined by MTT assay. (b) AGS cells were treated with 30 μM Hispidulin for the indicated times. Total ERK1/2, phosphor-ERK1/2 (pERK1/2), total p38, phosphor-p38 (pp38), total JNK, and phosphor-JNK (pJNK) were detected by Western blot. (c) Attenuation of Hispidulin-induced NAG-1 upregulation in AGS cells by ERK inhibitor PD98059. AGS cells where treatment with 25–50 μM of PD98059 and NAG-1 expression was evaluated by Western blot analysis. (d) Apoptosis induced by Hispidulin was attenuated by PD98059 ERK inhibitor treatment followed by flow cytometry analysis. Each column represents the mean ± SD (*P < 0.05; **P < 0.01; ***P < 0.001). Expression of β-actin was used as an internal control.

upregulation and growth inhibition, we applied MAPK inhibitors to Hispidulin-treated AGS cells. The ERK inhibitor PD98059 reduced the growth inhibitory effects of Hispidulin significantly (Figure 5(a)); however, the p38 inhibitor SB203580 and the JNK inhibitor SP600125 did not limit its growth inhibitory effects in AGS cells (data not shown). The ERK1/2 inhibitor PD98059 was the sole compound to reduce Hispidulin-induced upregulation of EGR-1 and NAG-1 protein expression (Figure 5(c)). These data support the concept that Hispidulin-induced ERK1/2 activity is critically involved in the regulation of EGR-1 and NAG-1 expression.

In an attempt to identify the signaling pathway through which PI3K/AKT/GSK3β is involved in receptor signal transduction through tyrosine kinase receptors for hispidulin, the effect of LY294002, a PI3K inhibitor, was examined. We found that LY294002 did not reverse tumor growth inhibition caused by hispidulin (data not shown). Compared with the effects of other phytochemical agents, this result suggests that the ability of our drug to cause tumor apoptosis might not go through this pathway. Results from analyses of COX-2 protein expression in AGS cells following their treatment with Hispidulin showed that Hispidulin induced morphological changes in the AGS cells (data not shown), inhibited AGS cell growth (Figure 1(a)), and arrested the cell cycle at the G1/S phase (Figure 3(a)). The tumor suppressor protein p53 plays a role in the molecular response to DNA damage. Acting as a DNA-binding transcription factor, it regulates specific target genes to arrest the cell cycle and initiate apoptosis. Following...
In summary, Hispidulin is a novel flavonoid compound that has modulatory effects on the expression of cyclin D1/cyclin E GI/S regulatory proteins and initiates the apoptotic cascade [19, 20]. Our study findings show that NAG-1 displays antitumorigenic and proapoptotic activities in vitro and indicate that Hispidulin, regulates NAG-1 expression. Although further detailed analyses are required to fully elucidate the mechanisms underlying the antitumorigenic effects of Hispidulin these results should encourage further investigation of Hispidulin as a potential novel clinical anticancer drug.

Abbreviations

NAG-1: NSAID-activated gene-1
K8: Isochaialulactone
RT-PCR: Reverse transcription-polymerase chain reaction
DMSO: Dimethyl sulfoxide
PBS: Phosphate-buffered saline
PCR: Polymerase chain reaction
MTT: 3-(4,5-dimethyl thizol-2-yl)-2,5-diphenyl tetrazolium bromide
PVDF: Polyvinyldenefluoride
MBD: Methyl-CpG binding domain
NSAIDs: Anti-inflammatory drugs
CI: Combination index
FBS: Fetal bovine serum
IHC: Immunohistochemical.

Conflict of Interests

All of the authors indicated no potential conflict of interests relevant to this paper.

Authors’ Contribution

Due-Chuan Chan and Yi-Lin Sophia Chen contributed equally to this work.

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