Emodin Alleviates Sodium Taurocholate–Induced Pancreatic Ductal Cell Damage by Inhibiting the S100A9/VNN1 Signaling Pathway

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Objectives: Because the pathogenesis of the disease is unclear, the treatment of patients with acute pancreatitis, especially severe acute pancreatitis, is still a major challenge for clinicians. Emodin is an anthraquinone compound extracted from rhubarb that can alleviate the damage to pancreatic ductal epithelial cells induced by adenosine triphosphate, but whether it has a similar protective effect on sodium taurocholate (STC)–stimulated pancreatic ductal cells and the underlying mechanism has not yet been reported.

Methods: A model of STC-induced HPDE6-C7 human pancreatic ductal epithelial cell injury was established, and then apoptosis and the levels of reactive oxygen species (ROS), glutathione, gamma-glutamylcysteine synthetase, and inflammatory cytokines were assessed in the presence or absence of emodin pretreatment. S100 calcium binding protein A9 (S100A9) and Vanin1 (VNN1) protein expression was also measured.

Results: Emodin significantly increased HPDE6-C7 cell viability, inhibited apoptosis and ROS release, and elevated glutathione levels and gamma-glutamylcysteine synthetase activity. Furthermore, emodin downregulated S100A9 and VNN1 protein expression and inhibited the production of inflammatory factors, such as interleukin (IL)-1β, IL-6, IL-8, and IL-18.

Conclusions: Emodin attenuates STC-induced pancreatic ductal cell injury possibly by inhibiting S100A9/VNN1-mediated ROS release. This finding provides evidence for the future development of emodin as a therapeutic agent.

Key Words: acute pancreatitis, ductal cells, emodin, S100 calcium binding protein A9, Vanin1

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A cute pancreatitis (AP), defined as the rapid onset of pancreatic inflammation, is the most common gastrointestinal disease leading to hospital admission worldwide. The annual incidence of AP ranges from 13 to 45 per 100,000 individuals in the United States, and the mortality rate of severe AP (comprising approximately 10% of all cases) remains remarkably high (~28%). The typical characteristics of AP are acinar cell injury and exocrine abnormalities, and sustained inflammation is responsible for local or systemic inflammatory response syndrome. Thus, most studies have focused on pancreatic acinar cell damage mediated by various stimuli and its underlying molecular mechanisms. The exocrine pancreas is composed of both acinar and ductal cells. These 2 types of exocrine cells interact closely to maintain the normal physiological function of the pancreas. Pancreatic acinar cell destruction is undoubtedly the terminal step in pancreatitis, but the duct plays a crucial role in maintaining the integrity of the pancreas; when this protective function is compromised, pancreatitis occurs. Under physiological conditions, the primary function of the pancreatic duct is to secrete ductal fluid and HCO3−, which together form a ductal mucosal barrier that protects the pancreas from noxious stimuli by blocking the backflow of bile and trypsin into the pancreas. Previous studies have shown that high concentrations of bile acids inhibit HCO3− secretion, thus leading to damage to the epithelial barrier. The ducts can no longer protect against bile acids, and loss of this protective effect may contribute to the progression of AP. Given the anatomy and biological functions of the pancreatic duct, the pancreatic ductal epithelium seems to be vulnerable to harmful stimuli, such as endoscopic retrograde pancreatography and gallstone passing. However, the mechanisms of pancreatic duct injury in AP pathogenesis have not been clarified, and further exploration is warranted.

S100 calcium binding protein A9 (S100A9), also known as myeloid-related protein-14, is one of the primary members of the S100 protein family and has been demonstrated to play an important role in the regulation of migration or adherence of inflammatory cells. Our previous study showed that deletion of S100A9 in the pancreatic duct ameliorates AP by targeting Vanin1 (VNN1)–mediated reactive oxygen species (ROS) release, which provides evidence for S100A9 as a potential therapeutic target for AP.

The Chinese herb rhubarb has been used alone or as a monochromatic herb in traditional Chinese medicine (TCM) formulas (eg, Qingyi decoction, Dachengqi decoction, and Yinchenhao decoction) to treat AP for many years. Emodin (1,3,8-trihydroxy-6-methylanthraquinone) is an anthraquinone derivative mainly extracted from rhubarb and has effects against AP in vitro and in vivo. Recent studies on the effect of emodin against AP have focused mainly on its ability to protect pancreatic acinar cells and suppress the inflammatory cascade. Interestingly, our previous research found that emodin also ameliorates the damage to pancreatic ductal epithelial cells induced by adenosine triphosphate in vitro. In this article, a model of sodium taurocholate (STC)–induced damage was first established to explore the protective effect of emodin
against pancreatic duct injury in vitro. Emodin may alleviate STC-induced pancreatic ductal epithelial cell damage by inhibiting apoptosis and inflammation by targeting the S100A9/VNN1 pathway.

MATERIALS AND METHODS

Reagents and Materials

High-glucose Dulbecco's modified Eagle's medium was obtained from HyClone Co (Logan, Utah). Fetal bovine serum, trypsin-Ethylen Diamine Tetraacetic Acid (0.05%), and phenol red were provided by Gibco Co, Ltd (Thermo Fisher Scientific, Cleveland, Ohio). Emodin and STC were purchased from Solarbio Science and Technology Co (Beijing, China). PrimeScript RT reagent and SYBR PremixEx Taq II (TliRNaseH Plus) were purchased from TaKaRa Biotechnology Co, Ltd (Dalian, China). RNAex Pro RNA reagent was obtained from Accurate Biology Co, Ltd (Changsha, China). A ROS Detection Assay Kit was purchased from AmyJet Scientific Inc Co, Ltd (Wuhan, China). An α-Amylase (α-AMS) Assay Kit and Gamma-Glutamylcysteine Synthetase (γ-GCS) Assay Kit were provided by the Nanjing Jiancheng Institute of Biotechnology (Nanjing, China). A Micro Reduced Glutathione (GSH) Assay Kit was provided by Solarbio Science and Technology Co (Beijing, China). Rabbit anti-S100A9 was purchased from Cell Signaling Technology, Inc (Beverly, Mass). Rabbit anti-VNN1 was purchased from the Proteintech Group (Chicago, Ill). Goat anti-rabbit IgG H&L (Horesradish Peroxidase) was obtained from Abcam (Cambridge, UK). An Annexin V-FITC/PI Apoptosis Detection Kit was provided by Dalian Meilun Biotechnology Co, Ltd (Dalian, China).

Cell Culture

The HPDE6-C7 human pancreatic duct cell line was obtained from American Type Culture Collection (Manassas, Va). The cells were cultured in high-glucose Dulbecco's modified Eagle medium supplemented with 10% fetal bovine serum in a humid environment containing 5% CO2 at 37°C.

Cell Viability Assay

A cell viability assay was used to determine the cytotoxicity of emodin and the optimum concentration and duration of emodin treatment. Emodin was dissolved in dimethyl sulfoxide before being added to cells; the final concentration of dimethyl sulfoxide was less than 0.1%. Briefly, HPDE6-C7 cells were plated in 96-well plates at a density of 1 × 105 cells/mL and incubated for 24 hours. Then, the cells were incubated with the different concentrations of emodin (180, 90, 45, 22.5, 11.25, and 5.625 μM) for 24 hours in 5% CO2 at 37°C. After that, the cells in each well were incubated with 10 μL of 5 mg/mL 3-(4,5)-dimethylthiazol-2-yl)-5-(z-y1)-3,5-di-phenyltetrazoliumromide for 3 to 4 hours. Finally, to dissolve the formazan crystals, the cells were incubated with 100 μL of dimethyl sulfoxide with shaking, and the absorbance at 490 nm was measured with a microplate reader (BioTek, Burlington, Vt).

HPDE6-C7 cells in the control group were cultured in regular medium, and those in the STC group were stimulated with STC at a dose of 1016 μM for 1 hour. The cells in the STC + emodin group were pretreated with 5, 6, 11.25, 22.5, or 45 μM emodin for 24 hours before being incubated with STC (1016 μM) for 1 hour to analyze the effects of different concentrations of emodin or were pretreated with 45 μM emodin for 3, 6, 12, 24, or 48 hours before being incubated with STC (1016 μM) for 1 hour to analyze the effects of emodin treatment for different durations. Cell viability was determined as described above. Morphological changes in the stimulated cells were visualized with a contrast microscope (Olympus Corp, Tokyo, Japan).

Quantitative Real-Time Polymerase Chain Reaction

HPDE6-C7 cells were seeded in 6-well plates. The cells in the STC group were treated with STC (1016 μM) for 1 hour, those in the emodin group were treated with 45 μM emodin for 24 hours, and those in the STC + emodin group were pretreated with emodin for 24 hours before being incubated with STC (1016 μM) for 1 hour. Total RNA was extracted from HPDE6-C7 cells subjected to different treatment using the TRIZol Universal Total RNA Extraction Kit following the manufacturer's instructions. To assess RNA integrity, a small amount of total RNA was diluted in RNase-free water, and the absorbance at 260, 280, and 320 nm was measured with a SimpliNano Ultra Micro Spectrophotometer (Biochrom, Cambridge, UK). One microgram of RNA was reverse-transcribed into cDNA with a PrimeScript RT reagent kit on a G-1000 polymerase chain reaction (PCR) system (Bioneer Technology, Hangzhou, China). Relative mRNA expression was determined using real-time PCR with SYBR Premix Ex TaqTMII (TliRNaseH Plus) on an ABI 7500 Real-Time PCR system (Applied Biosystems, Foster City, Calif). The following conditions were used: amplification at 95°C for 30 seconds, dematuration at 95°C for 5 seconds, and annealing at 60°C for 34 seconds for 40 cycles. The primer sequences are listed in Table 1. The expression levels of all genes were normalized to the expression of β-actin as an endogenous control, and the fold changes among the different groups were calculated by the 2-ΔΔCT method for quantitative analysis.

Flow Cytometry

The percentage of apoptotic cells was determined using an annexin V-FITC/PI apoptosis detection kit. HPDE6-C7 cells were incubated in 6-well plates in different culture media simultaneously. First, HPDE6-C7 cells were digested with 0.25% trypsin, collected in phosphate buffered saline (PBS) and pelleted by centrifugation at 1000 rpm for 5 minutes. Then, the cells collected in binding buffer were stained with 5 μL annexin V-FITC and 5 μL propidium iodide (PI). Finally, the cells were cultured at room temperature away from light for 15 minutes and analyzed by flow cytometry (BD LSFRFortessa Cell Analyzer, BD Biosciences, Franklin Lakes, NJ).

Reactive oxygen species generation was assessed using a ROS Detection Assay Kit following the manufacturer's instructions. HPDE6-C7 cells from different groups were collected, and the cell pellets were resuspended in culture medium containing 1 × ROS label. A single-cell suspension was prepared by gently pipetting the cells up and down and incubating them for 30 minutes in the dark. The treated cells were analyzed by flow cytometry (BD LSFRFortessa Cell Analyzer, BD Biosciences).

| TABLE 1. Primer Sequences Used in the Present Work |
| Gene | Primers (5'-3') | Forward | Reverse |
| --- | --- | --- | --- |
| β-actin | CTCGGGCAATGAGTCTCCTGTG | TCTTCTATTGTCGTTGGATCC | CTCCTGGCAGTTCCTG |
| IL-1β | ATGATGGCGTATACAGTGCAGCAA | GTCCGAGATTTCGAGCTTGC | TCTTCTATTGTCGTTGGATCC |
| IL-6 | AAGGCCCAGAGGCTGCAGGAGA | TCTTCTATTGTCGTTGGATCC | TCTTCTATTGTCGTTGGATCC |
| IL-8 | CACCTGTTGCAAACATGACTTCCAA | TCTTCTATTGTCGTTGGATCC | TCTTCTATTGTCGTTGGATCC |
| IL-18 | TCTTCTATTGACCAAGGAAATCCG | TCTTCTATTGACCAAGGAAATCCG | TCTTCTATTGACCAAGGAAATCCG |
Quantification of Total GSH Levels

Total GSH levels were measured with a Micro Reduced GSH Assay Kit according to the manufacturer's instructions. No less than 10⁶ cells from each group were harvested in 1× PBS and collected by centrifugation. Then the cells were suspended by 3 times the volume of protein precipitant to the cell pellet, and the cells underwent freezing-thawing 2 to 3 times. After centrifugation, the supernatant was obtained and placed in a 96-well plate. According to a standard calibration curve, total GSH concentrations were determined using the kinetic method. The absorbance was measured at 412 nm with a microplate reader.

Measurement of γ-GCS Levels

Levels of γ-GCS were measured according to the instructions of the γ-GCS assay kit. Total protein was extracted from cells with a protein extraction kit. Glutathione promoter was added to the protein samples, and the samples were incubated at 37°C for 6 minutes. Then, the optical density was measured at a wavelength of 636 nm using a microplate reader.

Animal Experiment

Wild-type male C57BL/6 (20–22 g) mice were provided by the Experimental Animal Center of Dalian Medical University. All animal care and experimental procedures were conducted in accordance with the institutional guidelines for the care and use of laboratory animals and were approved by the Animal Care and Use Committee of Dalian Medical University (no. AEE21019). The mice were anesthetized by isoflurane inhalation. The common hepatic duct was temporarily clamped at the liver hilum to prevent hepatic reflux after a midline incision was made following anesthesia, pancreatic duct was temporarily clamped at the liver hilum to prevent hepatic reflux after a midline incision was made following anesthesia, and after 6 and 12 hours. The mice in both the control and STC groups were administered an equivalent volume of 1% carboxymethylcellulose sodium at the same time point. The mice were killed 24 hours after AP model establishment, serum was obtained from apical blood samples, 1 part of the pancreas was stored in 4% paraformaldehyde solution, and the remaining part was stored at −80°C.

Amylase Assay

To evaluate the effects of emodin on AP mice, α-AMS levels in the serum were measured with α-AMS assay kits. Serum samples were diluted 1:100 in PBS, and 100 μL of the final diluted serum was added to substrate buffer in a centrifuge tube. After incubation for 7.5 minutes at 37°C, iodine working solution was added, and the samples were mixed thoroughly. Then, the absorbance at 660 nm was read using a microplate reader.

Hematoxylin-Eosin Staining

Pancreatic injury was assayed through hematoxylin-eosin (HE) staining. Parafomaldehyde-treated tissue samples were embedded in paraffin wax, and 4-μm tissue sections were cut, dewaxed, rehydrated, and stained with HE. Images of the tissue sections were obtained under a light microscope (Olympus Corp, Tokyo, Japan) at ×200 magnification.

Western Blotting

Total protein was isolated from HPDE6-C7 cells using a protein extraction kit, and the protein concentration was quantified with a Bicinchoninic Acid protein assay kit. Then, the proteins were separated by sodium dodecyl sulfate-polyacrylamide gel electrophoresis (10%–12%) and transferred onto Polyvinylidene Fluoride membranes (EMD Millipore, Merck KGaA, Darmstadt, Germany). The membranes were incubated overnight at 4°C with primary antibodies against S100A9 (diluted 1:1000 in Tris-buffered saline with Tween), VNN1 (diluted 1:500 in Tris-buffered saline with Tween), α-AMS, and β-actin, and the protein expression was normalized to that of β-actin. Finally, protein bands were visualized by enhanced Chemiluminescence using a Tanon-5200 Digital Gel Image Analysis System (Tanon Science and Technology Co, Ltd, Shanghai, China). Gel-Pro Analyzer 4.0 software (Media Cybernetics, Rockville, Md) was used for quantitative analysis.

FIGURE 1. Emodin alleviated STC-induced HPDE6-C7 cell injury. A, The maximum dose of emodin administered in vitro was 45 μM. N = 6; compared with 0 μM, **p < 0.01. B, Emodin had no effect on normal HPDE6-C7 cell morphology. C, The optimal duration of 45 μM emodin treatment in vitro was 24 hours. N = 6. **p < 0.01, STC vs control; ##p < 0.01, emodin + STC vs STC. D, Emodin (45 μM) reversed STC-induced HPDE6-C7 cell morphology damage. Data are expressed as mean ± SEM. Cells were collected 24 hours after STC/emodin treatment.
Data and Statistical Analysis

The data are presented as the mean ± standard error of the mean (SEM) of at least 3 separate experiments. GraphPad Prism 7.0 software (GraphPad Software, Inc, San Diego, Calif) was used for statistical analysis. One-way analysis of variance followed by Tukey post hoc test (according to homogeneity of variances: yes/no) was used to analyze differences between multiple independent groups. \( P < 0.05 \) or \( < 0.01 \) was considered statistically significant.
RESULTS

Emodin Alleviated STC-Induced HPDE6-C7 Cell Injury

To identify the dose of emodin that did not injure HPDE6-C7 cells, the cytotoxicity of emodin was determined by a cell proliferation assay. Considering the obvious cytotoxicity of 90 μM emodin to HPDE6-C7 cells (P < 0.01), the maximum concentration of emodin used in vitro in this study was 45 μM (Fig. 1A). HPDE6-C7 cells were visualized by light microscopy, and no apparent damage to cell morphology was observed in the control or 45 μM emodin treatment group (Fig. 1B). Compared with that of control HPDE6-C7 cells, the viability of HPDE6-C7 cells treated with STC was significantly reduced, but 5.625, 11.25, 22.5, and 45 μM emodin increased cell viability in a dose-dependent manner. Furthermore, use of a time gradient verified that the optimal pretreatment duration for 45 μM emodin was 24 hours (Fig. 1C). Analysis of cell morphology showed that STC-stimulated HPDE6-C7 cells were mainly round and bright and that STC increased the number of dead HPDE6-C7 cells and decreased the total HPDE6-C7 cell number, whereas emodin (45 μM) reversed STC-induced HPDE6-C7 cell damage (Fig. 1D).

Emodin Regulated the Expression of S100A9/VNN1 Signaling–Related Proteins in STC-Stimulated HPDE6-C7 Cells

As shown in Figure 2A, emodin had no impact on the protein levels of S100A9 and VNN1 in normal HPDE6-C7 cells. The protein expression levels of S100A9 and VNN1 were notably increased after the addition of STC, whereas emodin reversed the increase in the protein expression of S100A9 and VNN1 induced by STC, as shown in Figure 5B, suggesting that emodin alleviates STC-induced HPDE6-C7 cell injury by inhibiting the S100A9/VNN1 signaling pathway.

Emodin Inhibited Oxidative Stress in STC-Treated HPDE6-C7 Cells

The levels of ROS, GSH, and γ-GCS were not significantly changed in the emodin treatment group compared with the control group (Figs. 3A, B). The ROS levels in the STC group were significantly increased compared with those in the control group, and after the administration of emodin, ROS levels were obviously decreased (Fig. 3C). In contrast, GSH and γ-GCS levels were decreased by STC, but increased by emodin (Fig. 3D). These results indicated that inhibition of S100A9/VNN1 signaling by emodin can restore redox homeostasis in STC-stimulated HPDE6-C7 cells.

Emodin Reversed STC-Induced Apoptosis of HPDE6-C7 Cells

Apoptosis of HPDE6-C7 cells was analyzed by FITC Annexin V-PI double labeling and flow cytometry. Cell apoptosis was not affected by treatment with 45 μM emodin, as shown in Figure 4A. As shown in Figure 4B, apoptosis of HPDE6-C7 cells was promoted in the STC stimulation group compared with the control group, and emodin treatment inhibited the induction of apoptosis by STC. The relative apoptosis rate after emodin treatment was higher than that in the control group but lower than that in the STC stimulation group. These results indicated that inhibition of S100A9/VNN1 signaling by emodin may alleviate STC-induced HPDE6-C7 cell apoptosis.

Emodin Reduced Inflammatory Factor Release From STC-Stimulated HPDE6-C7 Cells

We explored the effects of emodin on the expression of inflammatory factors in STC-stimulated HPDE6-C7 cells. Compared with those in the control group, the mRNA levels of interleukin (IL)-1β, IL-6, and IL-8 in the emodin treatment group were significantly decreased, whereas there were no significant changes in IL-18 mRNA levels after emodin treatment (Fig. 5A). The mRNA levels of IL-1β, IL-6, IL-8, and IL-18 in the STC group were significantly higher than those in the control group; however,
the expression levels of these factors in the STC + emodin (45 μM) group were lower than those in the STC group (Fig. 5B). These results indicated that blockade of the S100A9/VNN1 pathway by emodin can reduce the massive release of inflammatory factors caused by STC.

Emodin Relieved STC-Induced AP in Mice

As shown in Figure 6A, compared with those in mice exposed to STC, serum α-AMS levels in mice treated with emodin were significantly decreased. The protein levels of S100A9 and VNN1 in the pancreas were also decreased in mice treated with emodin compared with the STC group (Fig. 6B). Emodin also alleviated pancreatic damage in mice treated with STC (Fig. 6C).
VNN1 in the STC group were obviously higher than those in the control group; however, the expression levels of these proteins in the emodin (45 μM) group were lower than those in the STC group (Fig. 6B). In addition, HE staining showed obvious pancreatic atrophy, including vacuolization, increased neutrophils, and a small amount of bleeding, in mice treated with STC; however, these deleterious changes were significantly ameliorated by emodin. Moreover, emodin had no effect on the pancreatic tissue of normal mice (Fig. 6C). These results indicated that emodin can alleviate the STC-induced AP in mice by repressing the S100A9/VNN1 pathway.

**DISCUSSION**

At present, because most studies on AP have focused on physiological changes in pancreatic acinar cells, efforts to identify the drugs for AP are focused mainly on acinar cell injury.24,26 Interestingly, pancreatic ductal cells, as one of the main components of the exocrine pancreas, also play an important role in maintaining the physiological function of the pancreas.27 Our previous study revealed that pancreatic ductal cells are the first to be exposed to the toxic stimuli related to pancreatitis, which indirectly lead to structural and functional impairment of pancreatic acinar cells.16 Although many previous studies have confirmed that TCM compounds, Chinese herbs, and natural products have effects against AP; the mechanisms of action of TCM compounds against AP is still unclear.17,20,24 The compositions of TCM compounds are complex and diverse; each Chinese herb has its own characteristics and different processing and compatibility methods lead to different curative effects.25 Much effort is needed to determine the drug compositions and pharmacological characteristics of TCM compounds, and such investigations are tedious. Based on previous clinical and experimental research, the efficacy of TCM formulas, such as Dachengji decoction and Qingyi decoction, in the treatment of AP has been confirmed.28,29 As the key ingredients of these 2 decoctions, the active components of rhubarb have been successfully isolated and identified.30 Emodin is one of the main active components of rhubarb. Our previous studies confirmed that emodin alleviates STC-induced pancreatic acinar cell injury and inflammation, as well as adenosine triphosphate-stimulated pancreatic duct cell injury.24,25 In this study, we established a model of STC-stimulated HPDE6-C7 cell injury to explore whether emodin has a protective effect against pancreatic duct injury in vitro and its potential mechanisms. The results showed that the viability and morphology of STC-stimulated HPDE6-C7 cells treated with 45 μM emodin for 24 hours were significantly improved.

The calcium-binding protein S100A9 is abundantly expressed in monocytes/macrophages and neutrophils, and it is also located in the cytoplasm and plasma membrane of pancreatic cells under normal conditions in the human gastrointestinal system.31 S100A9 is rapidly released upon necrosis or actively secreted by damaged cells and acts as a damage-associated molecular pattern in response to inflammatory stimuli.32 S100A9 selectively binds to Ca2+, Zn2+, and actin, as well as receptor for advanced glycation end products and toll-like receptor 4, with strong affinity and has intracellular and extracellular regulatory effects associated with the inflammatory response and tumorigenesis.33 Shen et al34 found that plasma S100A9 levels in patients with acute ischemic stroke can be used as biomarkers for the therapeutic evaluation of 4 TCM formulations (Buyang Huanwu decoction, Xuefu Zhuyu decoction, Tianma Gouteng decoction, and Shengyu decoction) before functional recovery can be observed. However, to date, there is no related research on the efficacy of rhubarb and its active component emodin in the treatment of AP via regulation of S100A9 expression. In our previous study based on isobaric tags for relative and absolute quantitation proteomics, we found that the expression of S100A9 in the pancreatic tissues of rats with AP was significantly increased.35 Mechanistically, S100A9, as a key regulator of STC-induced pancreatic duct cell injury, may activate the PYD domain-containing protein 3 inflammase and aggravate AP through VNN-mediated ROS release.16 VNN1 is a glycosylated phosphatidylinositol-anchored extracellular enzyme that can hydrolyze pantethine into pantethenic acid and cysteamine, thereby inhibiting γ-GCS enzyme activity and GSH activity. Previous studies have evaluated the function of VNN1 in oxidative stress, showing that upregulation of VNN1 expression can lead to activation of the proinflammatory response in the microenvironment by reducing antioxidant activity.36,37 In this work, we explored the function of S100A9 as a potential target of emodin in the treatment of AP for the first time. By measuring the protein expression of S100A9 and VNN1, we found that emodin can effectively inhibit the S100A9/VNN1 signaling pathway in STC-treated HPDE6-C7 cells and reduce the mass release of ROS. We also measured GSH and γ-GCS activity in HPDE6-C7 cells and found that GSH levels and γ-GCS enzyme activity were significantly increased by emodin-mediated inhibition of the S100A9/VNN1 pathway.

Generally, emodin can significantly inhibit the expression of S100A9 in HPDE6-C7 cells stimulated with STC, thus inhibiting the release of ROS mediated by VNN1, increasing antioxidant activity in injured duct cells, inhibiting the inflammatory cascade and ameliorating pancreatic duct injury in AP. Therefore, S100A9 may be an important target of emodin in the treatment of AP.

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