Schisandra chinensis regulates drug metabolizing enzymes and drug transporters via activation of Nrf2-mediated signaling pathway

Abstract: Drug metabolizing enzymes (DMEs) and drug transporters are regulated via epigenetic, transcriptional, posttranscriptional, and translational and posttranslational modifications. Phase I and II DMEs and drug transporters play an important role in the disposition and detoxification of a large number of endogenous and exogenous compounds. The nuclear factor (erythroid-derived 2)-like 2 (Nrf2) is a critical regulator of a variety of important cytoprotective genes that are involved in disposition and detoxification of xenobiotics. Schisandra chinensis (SC) is a commonly used traditional Chinese herbal medicine that has been primarily used to protect the liver because of its potent antioxidative and anti-inflammatory activities. SC can modulate some DMEs and drug transporters, but the underlying mechanisms are unclear. In this study, we aimed to explore the role of Nrf2 in the regulatory effect of SC extract (SCE) on selected DMEs and drug transporters in human hepatocellular liver carcinoma cell line (HepG2) cells. The results showed that SCE, schisandrin A, and schisandrin B significantly increased the expression of NAD(P)H: Nicotinamide Adenine Dinucleotide Phosphate-oxidase or:quinone oxidoreductase 1, heme oxygenase-1, glutamate–cysteine ligase, and glutathione S-transferase A4 at both transcriptional and posttranslational levels. Incubation of HepG2 cells with SCE resulted in a significant increase in the intracellular level of glutathione and total glutathione S-transferase content. SCE significantly elevated the messenger ribonucleic acid and protein levels of P-glycoprotein and multidrug resistance-associated protein 2 and 4, whereas the expression of organic anion transporting peptide 1A2 and 1B1 was significantly downregulated by SCE. Knockdown of Nrf2 by small interfering ribonucleic acid attenuated the regulatory effect of SCE on these DMEs and drug transporters. SCE significantly upregulated Nrf2 and promoted the translocation of Nrf2 from cytoplasm to the nuclei. Additionally, SCE significantly suppressed the expression of cytosolic Kelch-like ECH-associated protein 1 (the repressor of Nrf2) and remarkably increased Nrf2 stability in HepG2 cells. Taken together, our findings suggest that the hepatoprotective effects of SCE may be partially ascribed to the modulation of DMEs and drug transporters via Nrf2-mediated signaling pathway. SCE may alter the pharmacokinetics of other coadministered drugs that are substrates of these DMEs and transporters and thus cause unfavorable herb–drug interactions.

Keywords: Nrf2, Keap1, HepG2 cell, drug metabolizing enzyme, drug transporter, P-gp, MRP, OATP, Schisandra chinensis

Introduction

There is an increasing popularity of natural products including Chinese herbal medicines that are used for the management of various ailments and improvement of body health, although the clinical evidence for their application is poor or lacking. An estimated 25% of adults in developed countries and more than 80% of the population in developing countries consume herbal medicines for disease treatment. The 2002 National Health
Interview Survey of the US (n=5,456) indicated that 18.9% of adults used herbal supplements within the past 12 months and 57.3% (n=3,315) of these people used herbs to treat a specific health condition. Based on the 2002 National Health Interview Survey, an estimated ~38.2 million adults in the US took herbal products and supplements in 2002. In a study including 61,587 individuals aged 50–76 years in the US, one-third of the population stated that they used herbal supplements. However, there are limited data on the pharmacokinetics, mechanisms of action, and safety profiles for most Chinese herbal medicines. There are minimal regulatory requirements for safety and efficacy of herbal medicines in Western countries. Many herbal medicines and phytochemicals are traditionally used as organ protectants or chemopreventive agents, since they can induce antioxidant response and the expression of drug metabolizing enzymes (DMEs) and drug transporters that are responsible for the disposition and excretion of a large number of xenobiotics. On the other hand, many Chinese herbal medicines have been reported to act as inducers or inhibitors of important DMEs and drug transporters, and this has raised a safety concern due to potential harmful herb–drug interactions. However, the underlying mechanisms for herbal medicine-induced alterations in the expression of DMEs and drug transporters are not fully understood. A number of DMEs and drug transporters are regulated by a panel of ligand-activated transcriptional factors, namely nuclear receptors, including pregnane X receptor (PXR/NR1I2), constitutive androstane/activated receptor (CAR/NR1I3), farnesoid X receptor (FXR/NR1H4), liver X receptor-α (LXRα/NR1H3), and hepatocyte nuclear factor 4α (HNF4α/NR2A1). These nuclear receptors serve as xenobiotic sensors to upregulate important DMEs and drug transporters that participate in the metabolism and excretion of xenobiotics.

The nuclear factor (erythroid-derived 2)-like 2 (Nrf2) is a ubiquitous transcriptional factor encoded by the NFE2L2 gene in humans and plays a critical role in the maintenance of cellular homeostasis. Under physiological conditions, Nrf2 is localized in the cytosol and associated with its negative regulator, the Kelch-like ECH-associated protein 1 (Keap1, also known as the Nrf2 repressor). The Nrf2 protein, upon synthesis, is rapidly degraded by the 26S proteasome exclusively in the cytoplasm of cells, with a half-life of 15–20 minutes. Keap1 is crucial for the rapid turnover of Nrf2 and functions as an adaptor for Nrf2 ubiquitination at the lysine residues of the Neh2 domain by a Cullin-3-dependent ubiquitin–ligase complex. Upon exposure to xenobiotics, Nrf2 dissociates from Keap1 and translocates into the nucleus where it heterodimerizes with other leucine zipper proteins such as Maf or other partners and binds to a cis-acting deoxyribonucleic acid (DNA) regulatory element, namely the antioxidant response element (ARE) in the upstream promoter region, and initiates the transcription of a wide array of cytoprotective target genes that participate in xenobiotic disposition and clearance from the body and maintain essential cellular defensive functions against myriad toxic products, including metalloids, chemical carcinogens, and radiation. Nrf2 deficiency or inactivation of Nrf2-mediated pathway can lead to pathological changes such as abnormal cellular survival and apoptosis and prolonged inflammation recovery. Nrf2 knockout mice have showed significantly increased susceptibility to cancer, neurodegeneration, premature senescence, lung injury and fibrosis, and inflammation. Nrf2 also plays a key role in the regulation of many DMEs and drug transporters that contribute to the disposition, detoxification, and clearance of a large number of xenobiotics. Most DMEs and drug transporters have a critical role in the detoxification of xenobiotics and endogenous compounds. For example, NAD(P)H:quinone oxidoreductase 1 (NQO1), heme oxygenase-1 (HO-1), uridine 5'-diphosphoglucuronosyltransferase–glucuronosyltransferases, glutamate–cysteine ligase, modifier subunit (GCLM), and glutathione (GSH) S-transferases (GSTs) are important contributors to xenobiotic disposition and detoxification. Drug transporters such as P-glycoprotein (P-gp), multidrug resistance-associated proteins (MRPs/ABCCs), and organic anion transporting polypeptides (OATPs) are critical for drug distribution, disposition, and excretion. As such, pharmacological activation of Nrf2 has been widely advocated as a useful strategy for cancer prevention and treatment of several diseases. Induction or inhibition of DMEs and drug transporters may cause considerable alteration in the profiles of absorption, distribution, metabolism, and excretion of drugs. Therefore, it is important to identify the modulatory factors and signaling pathways involved in the regulation of DMEs and drug transporters.

**Schisandra chinensis** (SC, Wuweizi in Chinese) is a traditional Chinese herbal medicine that has been used for...
thousands of years. It has a wide range of pharmacological activities, including antioxidant, hepatoprotective, antibacterial, anticancer, cardioprotective, and anti-inflammatory effects.\textsuperscript{40} SC is widely and mainly used to protect the liver in clinical settings in Asian countries, including the People’s Republic of China, Japan, and Korea.\textsuperscript{40} SC extract (SCE) contains a mixture of various bioactive compounds, and the lignanoids are the main bioactive components, including schisandrin (Sch); pseudo Sch; SchA (ie, deoxyschisandrin), SchB, and SchC; γ-Sch; schisandrols A and B; Wuweizi ethers A and B; and schisanhenol.\textsuperscript{41,42} Among them, SchB is the most abundant dibenzocyclooctadiene lignanoid present in SCE. Previous preclinical studies have found that SCE and its major active lignanoids protected the liver in various liver-injured models, with different mechanisms involved in its hepatoprotective activities.\textsuperscript{43–59} These include antioxidation, inhibition of cytokine production, inhibition of apoptosis, activation of Nrf2, and other possible mechanisms.\textsuperscript{51–59} However, the mechanism is not fully understood, and the role of Nrf2 in the hepatoprotection of SCE is not elucidated. In this study, we aimed to examine the role of Nrf2 in the modulating effects of SCE on important DMEs and drug transporters in human hepatocellular liver carcinoma (HepG2) cells.

Materials and methods

Chemicals and reagents

SC was purchased from Daxiang Pharmaceuticals Inc. (Guangzhou, Guangdong, People’s Republic of China). SchA (purity =98.0\%, relative molecular mass =416.51 Dal) and SchB (purity =98.0\%, Mr =400.46 Dal) were bought from the National Institute for Food and Drug Control, Beijing, People’s Republic of China. tert-Butylhydroquinone (rBHQ, purity =97\%, used as a known Nrf2 activator\textsuperscript{60,61}), 3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide (MTT), cycloheximide (CHX, a known protein synthesis inhibitor), and dimethyl sulfoxide (DMSO) were obtained from Sigma-Aldrich Inc. (St Louis, MO, USA). Dulbecco’s Modified Eagle’s Medium (DMEM), fetal bovine serum (FBS), and 0.25% trypsin were purchased from Gibco Inc. (Grand Island, NY, USA). TRIzol reagent and real-time polymerase chain reaction (RT-PCR) kit were purchased from TaKaRa Inc. (Dalian, Liaoning, People’s Republic of China). Radio immunoprecipitation assay buffer (Beyotime Bio Inc., Shanghai, People’s Republic of China) and super signal-enhanced chemiluminescence detection system were purchased from Amersham Biosciences Inc. (Piscataway, NJ, USA). The bicinechonic acid kit for protein concentration measurement was bought from Beyotime Bio Inc. (Shanghai, People’s Republic of China). Primary antibodies against human HO-1 and GCLM were purchased from Epitomics Inc. (Burlingame, CA, USA), and primary antibodies against Nrf2, MRP1, MRP2, OATP1A2, GADPH and Keap1 were bought from Abcam Inc. (Cambridge, MA, USA). ARE–luciferase plasmids and pEZX–PG04 control plasmids were purchased from GeneCopoeia Inc. (Rockville, MD, USA).

Cell culture

The HepG2 cell line was obtained from American Tissue Collection Center (ATCC, Manassas, VA, USA). Cells were maintained in DMEM supplemented with 10% FBS, 100 U/mL penicillin, and 100 U/mL streptomycin at 37°C in a humidified 5% CO\textsubscript{2} atmosphere. Medium was refreshed every 2–3 days.

SCE preparation

To prepare SCE, the raw materials were dissolved in 95% ethanol at 70°C with fresh solvent changed after each 180 minutes three times, followed by macroporous resin (AB-8) purification using 75% ethanol as the eluting solvent. To quantify the contents of SchA and SchB in SCE, a quota of SCE (20 µL) was subject to high-performance liquid chromatography (HPLC, LC-2010C) (Simadzu, Kyoto, Japan) equipped with an SPD-100MAVP detector and a Discovery® C-18 HPLC column (4.6×25 mm, 5 µm, Penomenex Nucleosil, Torrance, CA, USA). The column was eluted using H\textsubscript{2}O:methanol (v/v, 30:70) at a flow rate of 1.0 mL/minute for 35 minutes. The absorbance at 254 nm was monitored. To examine the total content of lignanoids of SCE, a quota of SCE (60 µL) was examined at 570 nm on an ultra-violet (UV)-6100S scanning spectrophotometer (Shanghai Precision Instrument Co. Ltd, Shanghai, People’s Republic of China).

MTT assay

The effect of SCE on the viability of HepG2 cells was evaluated using the MTT assay. Cells were seeded into a 96-well plate (4×10\textsuperscript{3} cells/well) and treated with various concentrations of SCE, SchA, SchB, and rBHQ for 24 hours. Following the treatment, the cells were incubated with 10 µL (5 mg/mL) MTT for 4 hours at 37°C. The medium was removed and the formazan crystals produced by viable cells were dissolved in 150 µL DMSO. The absorbance was detected using a microplate reader (THERMO Multiskan FC, MIT, USA) at wavelength of 570/630 nm.
NQO1–ARE luciferase reporter assay
HepG2 cells in a 96-well plate were transfected with a mixture of 0.5 µg of ARE–luciferase plasmids, 0.05 µg of pEZX–PG04 control plasmids (both from GeneCopoeia Inc.), and Lipofectamine 2000 reagent (Invitrogen). After 6 hours, the transfection mixture was replaced with fresh complete DMEM. Cells were treated with various concentrations of SCE for 24 hours. The Renilla and Firefly luciferase activities were measured using the Dual Luciferase Assay System (GeneCopoeia Inc.) with a microplate reader (THERMO Multiskan FC).

Total ribonucleic acid isolation and RT-PCR analysis
Total ribonucleic acid (RNA) was extracted from HepG2 cells using RNAiso Plus reagent according to the manufacturer’s instruction. The concentration of total RNA was examined by spectroscope at 260/280 nm. Following the RNA extraction, 1 µg of total RNA was used to synthesize complementary deoxyribonucleic acid (cDNA). The cDNA was reverse transcribed using the Super Script III First-Strand Synthesis System (TaKaRa, Tokyo, Japan). The resultant cDNA was subject to RT-PCR analysis using Power SYBR Green PCR Master Mix (TaKaRa) in ABI7500 Real-Time PCR system (Applied Biosystems Inc., Grand Island, NY, USA). The sequences of primers are listed in Table 1.

Western blotting assay
Cells were lysed with the RIPA buffer containing the protease inhibitor cocktail. The protein concentration was measured by bicinchoninic acid assay, and protein samples were mixed with 4× Laemmli loading buffer containing dithiothreitol. The mixture was denatured at 95°C for 5 minutes. An equal amount of protein (20 µg) was separated by a criterion 8% Tris–HCl gel (Bio-Rad, Hercules, CA, USA), and subsequently transferred onto a polyvinylidene difluoride membrane using a Trans-Blot semidy transfer cell (Bio-Rad). The membrane was blocked with 5% skim milk for 2 hours at room temperature and then probed with specific primary antibodies at 4°C overnight. Following that, the membranes were incubated with horseradish peroxidase-conjugated secondary antibodies at room temperature for 2 hours. Subsequently, the blots were visualized using the enhanced chemiluminescence detection system (Amersham Pharmacia Biotech., Piscataway, NJ, USA).

RNA interference
RNA interference experiments were performed using a specific pool of oligonucleotides against human Nrf2 (On-Targetplus SMART pool human NFE2L2; RIBOBIO Inc., Guangzhou, People’s Republic of China). According to the manufacturer’s instructions, HepG2 cells were transfected with 25 nM Nrf2 small interfering RNA (siRNA) (siNrf2) using Lipofectamine 2000 (Invitrogen) for 24 hours. The negative control was performed by transfection with a non-targeting silencing pool (On-Targetplus siGAPDH; RIBOBIO Inc.) at the same concentration as siNrf2. The silencing efficiency was evaluated after 24 hours by analyzing the gene expression level of target of interest.

Measurement of intracellular GSH level and total GST content
HepG2 cells were seeded in a six-well plate at a density of 5.0×10⁵ cells/well. After 24 hours, cells were treated with serum-free medium containing different concentrations of drugs for 24 hours. Then, cells were washed with phosphate-buffered saline twice and harvested using the lysis buffer. Cell lysate

Table 1 Sequences of primers designed for real-time polymerase chain reaction analysis

<table>
<thead>
<tr>
<th>Gene</th>
<th>Forward primer</th>
<th>Reverse primer</th>
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<tr>
<td>Nrf2</td>
<td>ATAGCTGACCGCAGATATC</td>
<td>CATGCACAGTGAATGCTCT</td>
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<tr>
<td>HO-1</td>
<td>ACGTCTATGCGTCTGAG</td>
<td>TGTGGGGAAGTTGGAAGAG</td>
</tr>
<tr>
<td>NQO1</td>
<td>CAGTGTTGTGAATCCCTGCC</td>
<td>TCCCGTGGATCCTGCCAG</td>
</tr>
<tr>
<td>GCLM</td>
<td>AATCTTGCTCTGCTGAGTGA</td>
<td>TGGCTCTTGTGAGAATGCG</td>
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<tr>
<td>GSTA4</td>
<td>AGTGTGAGATGTAACACCC</td>
<td>ATGGCCTAAAGATGTGGAAGG</td>
</tr>
<tr>
<td>MRPI</td>
<td>CTCCGGGAATCAACGATCTGG</td>
<td>TAGAATTTGGTGTGTCACATTCT</td>
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<td>MRPI</td>
<td>GGATCCAAAGATGTAAGTATTTAAT</td>
<td>TCACAGTGCTGTCGAAATAG</td>
</tr>
<tr>
<td>Pgp</td>
<td>TGCTCAGACAGTGTGAGTTG</td>
<td>AATTCACGAAAGCTGGAAAC</td>
</tr>
<tr>
<td>OATP1A2</td>
<td>AAGCACAAGCGAGATCATTCA</td>
<td>GAGTTTCCACCATCAGCTTACATCA</td>
</tr>
<tr>
<td>OATP1B1</td>
<td>TGAAATCACCTGAGCTGTGTTT</td>
<td>CTCCTAGTGCAGTATACCATAGT</td>
</tr>
<tr>
<td>GAPDH</td>
<td>AGAAGGCTGGCCAGCTTGG</td>
<td>AGGGGCCATCCAGCTTGC</td>
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Abbreviations: Nrf2, nuclear factor (erythroid-derived 2)-like 2; HO-1, heme oxygenase-1; NQO1, NAD(P)H:quinone oxidoreductase 1; GCLM, glutamate–cysteine ligase, modifier subunit; GSTA4, glutathione S-transferase A4; MRPI, multidrug resistance-associated protein; Pgp, P-glycoprotein; OATP, organic anion transporting polypeptide; GAPDH, glyceraldehyde 3-phosphate dehydrogenase.
(20 µL) was used for protein concentration measurement and 6.5% (wt/vol) sulfosalicylic acid was added to samples at a ratio of 1:4 (v/v) for protein precipitation. Samples were incubated on ice for 10 minutes and then centrifuged at 1,000 x g for 10 minutes. The GSH level was immediately detected using a microplate reader at 405 nm, and the total GST content was measured at 412 nm.

**Statistical analysis**

Data are presented as the mean ± standard deviation from at least three independent experiments. Statistical analyses were performed by SPSS 16.0 software (SPSS Inc., Chicago, IL, USA) using one-way analysis of variance. P<0.05 is considered as statistical significance.

**Results**

**Determination of major components of SCE**

As is shown in Figure 1, both SchA and SchB were eluted by the mobile phase in the HPLC system. The retention time for SchA and SchB was 19.17 and 26.34 minutes, respectively. The linearity for SchA and SchB determination by HPLC was found in the range of 0.012−0.240 mg/mL (r=0.9999) and 0.0125–0.2500 mg/mL (r=0.9999), respectively. The content of SchA in SCE was 1.577 mg/g crude extracts and 0.763 mg/g crude extracts for SchB. A UV spectrophotometer was used to determine the total lignanoids of SCE with SchB used as the reference compound. The linearity was in the range of 4.84–24.20 mg/L (r=0.9948) and the total lignanoid content in SCE prepared was 39.77 mg/mL.

**Effect of SCE, SchA, and SchB on HepG2 cell viability**

First, in order to evaluate the cytotoxic effect of SCE and its major active lignanoids on HepG2 cells, the cell viability was examined using the MTT assay. Cells were incubated with SCE, SchA, SchB, and tBHQ for 24 hours, which resulted in a decrease in cell viability in a concentration-dependent manner (Figure 2). The cell viability was 96.1%±10.3%, 75.8%±10.3%, 63.4%±2.4%, 53.1%±3.9%, 28.4%±14.4%, and 15.25%±7.0% when HepG2 cells were treated with SCE at 15, 30, 60, 120, 240, and 480 µg/mL for 24 hours, respectively (Figure 2A). The cell viability was 93.3%±17.4%, 83.1%±19.7%, 72.3%±2.5%, 62.0%±2.5%, 40.3%±17.1%, and 28.3%±9.7% with the incubation of SchA at 3.125, 6.25, 12.5, 19.168 µM, and 26.335 µM, respectively. The effect of tBHQ, a known nuclear factor (erythroid-derived 2)-like 2 activator, at 3.125−100 µM on cell viability, was also examined (Figure 2B). Cell viability was evaluated by 3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide assay. The data shown are the mean ± standard deviation of at least three independent experiments.

**Abbreviations:** Sch, schisandrin; tBHQ, tert-butylhydroquinone; IC_{50}, half maximal inhibitory concentration.
25, 50, and 100 µM for 24 hours, respectively (Figure 2B). After cells were treated with SchB at 12.5, 25, 50, 100, 200, and 400 µM, the cell viability was 98.4%±27.2%, 82.7%±6.2%, 83.5%±27.7%, 72.7%±34.1%, 45.8%±22.9%, and 30.2%±10.8%, respectively. When the cells were treated with rBHQ at 3.125, 6.25, 12.5, 25, 50, and 100 µM, the cell viability was 97.0%±11.8%, 87.3%±36.0%, 74.9%±11.0%, 62.1%±24.6%, 45.3%±11.1%, and 16.5%±10.0%, respectively (Figure 2B). The half maximal inhibitory concentration values for SCE, SchA, SchB, and rBHQ were 306.1 µg/mL, 70.6 µM, 266.4 µM, and 50.7 µM, respectively.

These compounds only showed a low cytotoxicity toward HepG2 cells.

**SCE, SchA, and SchB induce the expression of selected DMEs in HepG2 cells**

SCE has been reported to be a potent antioxidant that is primarily used to treat liver diseases.40 Herein, the effect of SCE on the expression of several important DMEs was examined at transcriptional and posttranscriptional levels. HepG2 cells were incubated with SCE at 100, 150, 200, 250, and 300 µg/mL for 24 hours; the gene expression level of \( NQO1 \) was increased 1.9-, 2.6-, 1.6-, 2.5-, and 3.2-fold, respectively (\( P \leq 0.05 \)) (Figure 3A); and the gene expression level of GCLM was elevated 1.4-, 4.2-, 1.4-, 2.2-, and 3.2-fold, respectively (Figure 3B). The gene expression level of HO-1 was increased 1.4-, 2.6-, 1.6-, 2.5-, and 3.2-fold, respectively (\( P \leq 0.05 \)) (Figure 3C); and the gene expression level of GSTA4 was elevated 1.4-, 2.6-, 1.6-, 2.5-, and 3.2-fold, respectively (\( P \leq 0.05 \)) (Figure 3D).

**Figure 3** Effect of Schisandra chinensis extract (SCE), schisandrin (Sch) A, and SchB on the expression of NAD(P)H:quinone oxidoreductase 1 (\( NQO1 \)), glutamate–cysteine ligase, modifier subunit (GCLM), heme oxygenase-1 (HO-1), and glutathione S-transferase A4 (GSTA4) in human hepatocellular liver carcinoma cell line cells. Human hepatocellular liver carcinoma cell line cells were treated with SCE at 100–300 µg/mL, 20 µM SchA, or 200 µM SchB for 24 hours. Total ribonucleic acid was extracted from the treated cells and then the first-strand complementary deoxyribonucleic acid was synthesized. A quota of complementary deoxyribonucleic acid (2 µL) was used as the template for quantitative polymerase chain reaction analysis of the target genes in each sample. (A–D) Bar graphs show the messenger ribonucleic acid expression levels of \( NQO1 \), GCLM, HO-1, and GSTA4, respectively. Data are presented as the mean ± standard deviation (n=3).

**Notes:** *\( P \leq 0.05 \); **\( P \leq 0.01 \); ***\( P \leq 0.001 \) by one-way analysis of variance; Ctrl, control.
3.5-fold, respectively ($P<0.05$) (Figure 3B). Treatment of HepG2 cells with 250 and 300 $\mu$g/mL SCE resulted in a 2.1- and 3.9-, 1.5- and 1.3-fold increase ($P<0.01$) in the gene expression level of HO-1 (Figure 3C) and GSTA4 (Figure 3D), respectively. Moreover, there was a significant increase in the gene expression of NQO1, GCLM, and HO-1 when cells were incubated with SchA at 20 $\mu$M for 24 hours. Treating cells with 200 $\mu$M SchB also significantly induced the gene expression of NQO1 and GSTA4. On the other hand, the protein expression level of HO-1 and GCLM was also examined in order to evaluate the effect of SCE on the post-transcriptional regulation of DMEs. Incubation of HepG2 cells with SCE at 100, 150, 200, 250, and 300 $\mu$g/mL for 24 hours concentration dependently increased the expression levels of HO-1 and GCLM (Figure 4). SchA at 20 $\mu$M and SchB at 200 $\mu$M also significantly increased the expression levels of HO-1 and GCLM in HepG2 cells (Figure 4).

**SCE and SchB differentially modulate the expression levels of selected drug transporters in HepG2 cells**

Drug transporters play an important role in the excretion of xenobiotics and thus protect important organs from their toxicities. On the other hand, the regulatory effect of xenobiotics, including drugs and herbal medicines, on drug transporters may have substantial impact on the pharmacokinetic and pharmacodynamic profiles of a coadministered drug. As such, the effect of SCE on selected important drug transporters was evaluated at both transcriptional and post-transcriptional levels. Incubation of HepG2 cells with SCE significantly increased the gene expression level of P-gp, ABCB1, ABCG2, and ABCA1.

**Figure 4** Effect of SCE, SchA, and SchB on the expression of HO-1 and GCLM in human hepatocellular liver carcinoma cell line cells. Human hepatocellular liver carcinoma cell line cells were treated with SCE at 100–300 $\mu$g/mL, 20 $\mu$M SchA, or 200 $\mu$M SchB for 24 hours. Cellular lysates were subjected to Western blotting assay. (A) Representative blots of HO-1 and GCLM. (B, C) Bar graphs to show the expression levels of HO-1 and GCLM, respectively. Nicotinamide adenine dinucleotide phosphate was used as the internal control. Data are presented as the mean ± standard deviation from at least three independent experiments.

**Notes:** **$P<0.001$; ***$P<0.001$ by one-way analysis of variance.

**Abbreviations:** SCE, Schisandra chinensis extract; Sch, schisandrin; HO-1, heme oxygenase-1; GCLM, glutamate–cysteine ligase, modifier subunit; GAPDH, glyceraldehyde 3-phosphate dehydrogenase; Ctrl, control.
Figure 5 Effect of Schisandra chinensis extract (SCE), schisandrin (Sch) A, and SchB on the messenger ribonucleic acid expression of P-glycoprotein (P-gp), multidrug resistance-associated protein (MRP) 2, MRP4, organic anion transporting peptide (OATP) 1A2, and OATP1B1 in human hepatocellular liver carcinoma cell line cells. Human hepatocellular liver carcinoma cell line cells were treated with SCE at 100–300 µg/mL, 20 µM SchA, or 200 µM SchB for 24 hours. Total ribonucleic acid was extracted and complementary deoxyribonucleic acid was synthesized. A quota of complementary deoxyribonucleic acid (2 µL) was used for real-time polymerase chain reaction analysis of the target genes. (A–E) Bar graphs to show the relative expression levels of P-gp, MRP2, MRP4, OATP1A2, and OATP1B1, respectively. tert-Butylhydroquinone (tBHQ) was used as a known nuclear factor (erythroid-derived 2)-like 2 activator. Data are presented as the mean ± standard deviation (n=3).

Notes: *P<0.05; **P<0.01; ***P<0.001 by one-way analysis of variance; Ctrl, control.
MRP2, and MRP4 (Figure 5). There was a 2.0- and 1.5-fold increase in the gene expression level of P-gp in HepG2 cells treated with 200 or 250 µg/mL of SCE, respectively (Figure 5C). Treatment of HepG2 cells with SCE at 100, 150, 200, 250, and 300 µg/mL increased 1.5-, 1.8-, 1.8-, 2.5-, and 2.0-fold in MRP2 gene expression level (Figure 5A) and elevated 1.9-, 1.8-, 2.2-, and 1.7-fold in MRP4 gene expression level, respectively (Figure 5B). SchB at 200 µM significantly increased the gene expression level of MRP2 and MRP4. In addition, tBHQ (a known Nrf2 agonist/activator) significantly increased the gene expression level of P-gp and MRP2. There was no significant effect of SchA at 20 µM on the gene expression level of P-gp, MRP2, and MRP4 (Figure 5).

On the contrary, incubation of HepG2 cells with SCE significantly reduced the gene expression level of OATP1A2 and OATP1B1. There was a 30.9% and 53.1% reduction in the gene expression of OATP1A2 when cells were treated with SCE at 200 and 300 µg/mL, respectively (Figure 5D), and there was a 41.6%, 42.8%, and 86.4% decrease in the gene expression of OATP1B1 when cells were incubated with SCE at 100, 150, and 250 µg/mL, respectively (Figure 5E). However, SchB at 200 µM increased 1.4-fold in the gene expression level of OATP1A2 (P<0.05). There was no significant effect of SchA at 20 µM on the gene expression level of OATP1A2 and OATP1B1 (Figure 5). Notably, incubation of HepG2 cells with tBHQ significantly decreased 37.9% and 27.3% of the messenger RNA (mRNA) levels of OATP1A2 and OATP1B1, respectively (Figure 5).

In addition, the protein expression levels of P-gp, MRP2, and OATP1A2 were examined in HepG2 cells treated with SCE, SchA, or SchB. As shown in Figure 6, incubation of HepG2 cells with 250 and 200 µg/mL SCE significantly increased 1.9- and 1.4-fold in the levels of P-gp and MRP2, respectively, whereas there was an 85.7% reduction in OATP1A2 level at a concentration of 250 µg/mL SCE. Treatment of cells with 10 µM tBHQ increased 2.3-fold in the expression level of MRP2. In addition, SchA at 20 µM significantly enhanced the expression of P-gp and MRP2 but decreased the expression level of OATP1A2. SchB at 200 µM significantly increased the expression level of P-gp but reduced the expression of MRP2 and OATP1A2 (Figure 6). These findings demonstrate that SCE, SchA, and SchB at different concentrations have distinct effects on the expression of various drug transporters; SchA and SchB are the major modulators of these drug transporters; and Nrf2 appears to play an important role in the regulation of these drug transporters.

SCE, SchA, and SchB increase intracellular GSH levels and total GST contents
Owing to their potent antioxidant activity, GSH and GSTs have an important role in maintaining essential cellular function against the cytotoxic effects of xenobiotics. Herein, the effect of SCE on intracellular levels of GSH and total GST content was examined in HepG2 cells. SCE remarkably elevated the intracellular level of GSH and the total content of GSTs in HepG2 cells. There was a 2.6-, 8.2-, 6.3-, and 4.0-fold increase in the intracellular level of GSH when HepG2 cells were treated with 150, 200, 250, and 300 µg/mL SCE, respectively (Figure 7A). Similarly, there was a 2.7-, 4.9-, 3.6-, and 2.5-fold increase in the total content of GSTs when cells were incubated with 150, 200, 250, and 300 µg/mL SCE, respectively (Figure 7B). SchA at 20 µM and SchB at 200 µM showed a comparable promoting effect on the intracellular level of GSH and the total content of GSTs as SCE in HepG2 cells.

SCE activates Nrf2 in HepG2 cells as indicated by the NQO1–ARE luciferase assay
Following the observations of the regulatory effect of SCE on selected DMEs, drug transporters, and redox status in HepG2 cells, we further investigated the role of Nrf2 in the regulatory effect of SCE and its major components SchA and SchB on DMEs and drug transporters. It has been recognized that Nrf2 is a critical nuclear transcriptional factor regulating the expression of Phase II cytoprotective genes in response to xenobiotic exposure and other external stimuli.20,22 Herein, we examined the effect of SCE on the expression/activity of Nrf2 by testing the activity of NQO1–ARE luciferase, which indicates the activity of Nrf2. Incubation of HepG2 cells with SCE significantly enhanced the activity of Nrf2. There was a 2.2-, 3-, 6-, 6.1-, and 3.2-fold increase in the level of NQO1–ARE luciferase when cells were treated with SCE at 50, 100, 150, 200, 250, and 300 µg/mL, respectively. tBHQ at 10 µM, SchA at 20 µM, and SchB at 200 µM increased 4.4-, 4.0-, and 4.3-fold in the level of NQO1–ARE luciferin, respectively (P<0.01) (Figure 8). These results suggest that the SCE acts as an Nrf2 activator in HepG2 cells.

Nrf2 knockdown attenuates the induction of DMEs and efflux drug transporters
Nrf2 is a key transcriptional factor that regulates DMEs and drug transporters.20,22 To clarify the role of Nrf2 in the
Figure 6 Effect of SCE, SchA, and SchB on the protein expression of P-gp, MRP2, and OATP1A2 in human hepatocellular liver carcinoma cell line cells. Human hepatocellular liver carcinoma cell line cells were treated with SCE at 100–300 μg/mL, 20 μM SchA, or 200 μM SchB for 24 hours. Cellular lysates were subjected to Western blotting assay. (A) Representative blots of P-gp, MRP2, and OATP1A2. (B–D) Bar graphs to show the expression levels of P-gp, MRP2, and OATP1A2, respectively. Nicotinamide adenine dinucleotide phosphate was used as the internal control. Data are presented as the mean ± standard deviation from at least three independent experiments.

Notes: *P < 0.05; **P < 0.01; ***P < 0.001 by one-way analysis of variance.

Abbreviations: SCE, Schisandra chinensis extract; Sch, schisandrin; P-gp, P-glycoprotein; MRP2, multidrug resistance-associated protein 2; OATP1A2, organic anion transporting peptide; tBHQ, tert-butylhydroquinone; GAPDH, glyceraldehyde 3-phosphate dehydrogenase; Ctrl, control.

regulation of DMEs and drug transporters by SCE, RNA interference was performed to knock down Nrf2 in HepG2 cells. In comparison with the control group (si-Ncontrol), incubation of HepG2 cells with Nrf2 siRNA significantly decreased 50% and 48% in the gene expression of NQO1 (Figure 9C) and MRP2 (Figure 9E), respectively. There was a 30% reduction in the gene expression of HO-1 when treated with Nrf2 siRNA (P < 0.05) (Figure 9D). Furthermore, there was a 60% decrease in the expression level of Nrf2 compared with the control, and the inducing effect of SCE on the expression of Nrf2 was attenuated (Figure 9A and B). These results indicate that Nrf2-mediated signaling pathway is involved in the regulatory effect of SCE on DMEs and drug transporters in HepG2 cells.

SCE, SchA, and SchB suppress Keap1 expression and facilitate nuclear translocation of Nrf2 in HepG2 cells

Keap1 is known as an Nrf2 repressor that binds Nrf2 with cellular actin cytoskeleton in cytoplasm.20–22 We have observed that Nrf2 plays an important role in the regulatory effect of SCE on DMEs and drug transporters. Next, we further examined the effect of SCE, SchA, and SchB on Keap1 expression and the nuclear translocation of Nrf2 to clarify the regulatory effect of SCE on the interaction between Keap1 and Nrf2. Incubation of HepG2 cells with SCE concentration dependently decreased the cytosolic level of Keap1 (P < 0.01) (Figure 10). Treatment of HepG2 cells with SCE at 250
and 300 µg/mL significantly decreased the cytosolic level of Keap1 by 45.85% and 70.68%, respectively (P<0.001). SchA at 20 µM and SchB at 200 µM significantly decreased the cytosolic level of Keap1 by 32.94% and 35.90%, respectively (P<0.05). We also examined the effect of SCE on Nrf2 protein expression in the nucleus and cytoplasm. The results showed an increase in Nrf2 at both total and nuclear protein levels, which indicates that SCE facilitates the nuclear translocation of Nrf2 (Figure 10).

**SCE increases Nrf2 protein stability**

In addition, to further elucidate how SCE regulated DMEs and drug transporters through Nrf2 activation, the effect of SCE on Nrf2 protein stability was examined. HepG2 cells were treated with the protein synthesis inhibitor CHX at indicated times with or without pretreatment of SCE, SchA, and SchB. At the basal level, the half-life of Nrf2 protein was 13.07±0.50 minutes. Pretreatment of cells with SCE at 200 µg/mL, SchA at 20 µM, and SchB at 200 µM significantly prolonged the half-lives of Nrf2 protein (P<0.001), which were 26.02±0.46, 24.14±0.80, and 19.41±1.96 minutes, respectively (Figure 11). These results indicate that SCE, SchA, and SchB remarkably increase Nrf2 stability primarily through inhibition of protein degradation. Since SCE, SchA, and SchB downregulated Keap1, this would contribute to the prolonged half-lives of Nrf2 in HepG2 cells.

**Discussion**

SCE (Wuweizi), dried mature fruit from SC, is one of the most commonly used Chinese herbal medicines as a liver protectant that has been documented in *Sheng Nong’s Herbal Classic*. To date, it has been shown that SCE has remarkable protective effects on the liver, digestive system, central nervous system, and cardiovascular system. These beneficial effects can be ascribed to its antioxidative and...
Figure 9 Effect of Nrf2 knockdown by ribonucleic acid interference on the expression of important drug metabolizing enzymes and drug transporters. (A) Transfection efficiency of various Nrf2–small interfering (si) ribonucleic acids in human hepatocellular carcinoma cell line cells. (B–E) Bar graphs to show the gene expression levels of Nrf2, NQO1, HO-1, and MRP2, respectively. Data are presented as the mean ± standard deviation (n=4).

Notes: *P<0.05; **P<0.01; ***P<0.001 by one-way analysis of variance.

Abbreviations: Nrf2, nuclear factor (erythroid-derived 2)-like 2; NQO1, NAD(P)H:quinone oxidoreductase 1; HO-1, heme oxygenase-1; MRP2, multidrug resistance-associated protein; GAPDH, glyceraldehyde 3-phosphate dehydrogenase; SCE, Schisandra chinensis extract; tBHQ, tert-butylhydroquinone; Sch, schisandrin; Ctrl, control.
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SCE regulates DMEs and transporters through Nrf2 activation

anti-inflammatory activities. In the present study, we have observed that SCE upregulates important DMEs and drug transporters that participate in xenobiotic detoxification with the involvement of Nrf2-mediated signaling pathway in HeG2 cells.

The choice of SCE concentrations (150−300 µg/mL) in this study is based on the dosage of SCE used in patients who are administered 1.0−6.0 g of SCE per day. This equals 20.0−120.0 µg/mL of SCE, assuming a standard adult contains 50 kg of water and SCE is evenly dissolved in the body’s water. Our prepared SCE at 200 µg/mL contains 0.76 µM SchA and 0.39 µM SchB. The concentrations of SchA and SchB used in this study were significantly higher than those in SCE but did not elicit remarkable cytotoxicity toward HepG2 cells. Notably, we tested the effects of low concentrations of SchA and SchB on the expression of DMEs and drug transporters in HepG2 cells in our pilot experiment, but we did not see any effects.

The liver is the primary organ in the biotransformation and detoxification of endogenous and exogenous compounds by a large number of Phase I and II DMEs. Of great importance, Phase II conjugating DMEs have an important role in the detoxification of drugs and environmental toxicants and the maintenance of liver function. Recently, studies have shown that Phase I and II detoxifying and antioxidant enzymes such as GSTs, GCLM, NQO1, and HO-1 are critical in the maintenance of redox balance in the liver. NQO1 is a flavin adenine dinucleotide-binding cytosolic protein that forms homodimers and reduces quinones to hydroquinones, inhibiting the production of radical species. NQO1 also has anti-inflammatory effects, scavenges superoxide anion radicals, and stabilizes p53 and other tumor suppressors. Mutations in the NQO1 gene have been associated with tardive dyskinesia, an increased risk of hematotoxicity after exposure to benzene, and susceptibility to cancer, Alzheimer’s disease, and cardiovascular diseases. HO-1 cleaves the heme ring at the α-methylene bridge to form either biliverdin or verdoglobin when the heme is still attached to a globin, with the release of CO and free iron. Biliverdin is subsequently converted to bilirubin by biliverdin reductase. Unlike HO-2, HO-1 is

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Figure 10 Effect of SCE, SchA, and SchB on the expression of nuclear and total Nrf2 and Keap1. Human hepatocellular liver carcinoma cell line cells were treated with SCE at 100−300 µg/mL, 20 µM SchA, or 200 µM SchB for 24 hours. Whole cellular and nucleic lysates were subject to Western blotting analysis. (A) Representative blots of nuclear and total Nrf2 and Keap1 in human hepatocellular liver carcinoma cell line cells. (B–D) Bar graphs to show the relative expression level of nuclear and total Nrf2 (B and C) and Keap1 (D), respectively. Nicotinamide adenine dinucleotide phosphate was used as the internal control. Data are presented as the mean ± standard deviation (n=3).

Notes: *P<0.05; **P<0.01; and ***P<0.001 by one-way analysis of variance.

Abbreviations: SCE, Schisandra chinensis extract; Sch, schisandrin; Nrf2, nuclear factor (erythroid-derived 2)-like 2; Keap1, Kelch-like ECH-associated protein 1; GAPDH, glyceraldehyde 3-phosphate dehydrogenase; Ctrl, control.
an inducible isoform in response to stress such as oxidative stress, hypoxia, heavy metals, and cytokines. GCL is the first rate-limiting enzyme of GSH synthesis.70 Human GSTs contribute to the GSH conjugation or GSH-dependent biotransformation of xenobiotics, and many catalyze GSH peroxidase or thiol transferase reactions. GSTs also catalyze GSH-dependent isomerization reactions required for the synthesis of several prostaglandins and steroid hormones and the catabolism of tyrosine. Furthermore, GSTs serve as significant regulators of pathways determining cell proliferation and survival and as regulators of ryanodine receptors that are essential for muscle function.71 Dysregulation of NQO1, GCLM, GSTs, and HO-1 results in disruption of redox homeostasis in the liver or other organs when the level of reactive oxygen species outweighs the antioxidants under oxidative stress.72 Consequently, disrupted redox homeostasis may cause liver and organ damage. In the present study, we have observed that SCE significantly induced the expression of NQO1, GCLM, GSTA4, and HO-1 at both transcriptional and translational levels in HepG2 cells. These results can, at least in part, explain the antioxidative and hepatoprotective effects of SCE.

GSH is a ubiquitous intracellular tripeptide with diverse functions including detoxification, antioxidant defense, maintenance of thiol status, and modulation of cell proliferation.73 GSH is synthesized in the cytosol of mammalian cells in a tightly regulated manner. The major determinants of GSH synthesis are the availability of cysteine, the sulfur amino acid precursor, and the activity of the rate-limiting enzyme GCL, which is composed of a catalytic (GCLC) and modifier (GCLM) subunit.73 GSH has an important role in maintaining liver function via the regulation of redox homeostasis, and inadequate GSH levels have a significant association with toxicant-induced hepatotoxicity.74 Restoration of the ratio of GSH to its oxidized form can ameliorateoxicant-induced hepatotoxicity. In our study, we found that incubation of HepG2 cells with SCE remarkably elevated the intercellular level of GSH, which could partially explain the hepatoprotective effect of SCE. Moreover, we also observed that SCE significantly increased the total GST content that plays a critical role in the detoxification of a large number of xenobiotics. Taken together, the antioxidative effect of SCE can be ascribed, at least in part, to the upregulation of antioxidative enzymes and the elevation of intracellular levels of GSH. Given the effects of SCE on GSH and GSTs, SCE shows a coordinated regulatory effect in response to oxidative stress or chemical stimuli to exert its liver protective effects.

Drug transporters have an important role in cellular protection because of their capability of removing xenobiotics and endogenous substrates that can accumulate in cells and lead to toxicity.74,75 The ATP-binding cassette (ABC) transporters, such as P-gp (MDR1), MRP2, and MRP4, are responsible for the efflux of a number of endogenous and exogenous compounds.76–78 ABC transporters are divided into seven distinct subfamilies, including ABC1, MDR/TAP, MRP, ALD, OABP, GCN20, and white. P-gp is located on the apical surface of intestinal epithelial cells, bile canaliculi, renal tubular cells, and placenta and the luminal surface of capillary endothelial cells in the brain and testes. MRP2 is generally localized at the apical membrane, and MRP4 is at the sinusoidal membrane of hepatocytes.76–78 P-gp transports various substrates across the cell membrane, including
In the intestine, OATP1A2 is localized to the brush border of the OATP/SLCO21 family that belongs to uptake and SchB are inhibitors for P-gp and MRP1. SchB are substrates of P-gp. In the present study, we found that SCE upregulated the expression of OATP1A2 and OATP1B1 at transcriptional and posttranscriptional levels. In contrast to SCE, SchB at 200 µM upregulated the expression of OATP1A2. Nrf2 may contribute to this effect of SchB, but the involvement of other regulators and related signaling pathways such as FXR and protein kinase C cannot be excluded.

In our study, we further revealed the role of Nrf2-mediated signaling pathway in the regulatory effect of SCE and its main components on DMEs and drug transporters. This is consistent with the reports from several previously published studies where Nrf2 was found to play a role in the antioxidative and anti-inflammatory effects of SCE and its main active compounds. Activation of Nrf2-mediated pathway by pharmacologic approach serves to stimulate coordinated induction of hepatic DMEs such as GSTs and efflux transporters such as P-gp, MRP2, MRP3, and MRP4 upon binding to the ARE at the proximal region of the promoter in the target genes.

Nrf2 signaling is known as a mastery pathway that regulates the expression of a number of DMEs and transporters. Besides, Nrf2-mediated signaling pathway is widely involved in cell protection in cancer chemoprevention and anti-inflammatory process. Nrf2 belongs to the basic Lucien zipper nuclear transcriptional factor family, which shares regions of homology with the Cap ‘n’ Collar (CNC) structure. Nrf2 contains six highly conserved domains known as Nrf2–ECH homology (Neh) domains.

The Neh1 domain is a CNC–bZIP domain that allows Nrf2 to heterodimerize with small Maf proteins; the Neh2 domain allows for binding of Nrf2 to its cytosolic repressor Keap1.
the Neh3 domain plays a role in Nrf2 protein stability and may act as a transactivation domain, interacting with components of the transcriptional apparatus; the Neh4 and Neh5 domains also act as transactivation domains but bind to a different protein called cAMP response element binding protein, which possesses intrinsic histone acetyltransferase activity; and the Neh6 domain may contain a degron that is involved in the degradation of Nrf2.\textsuperscript{35,44,45} Nrf2 is considered to play an important role in the cellular defense mechanisms against xenobiotic toxicity and carcinogenicity.\textsuperscript{20,22} Activation of Nrf2 signaling pathway can increase the expression of the rate-limiting enzymes in GSH synthesis, thus enhancing cells’ capability against oxidative stress. It is evident that many small natural or synthetic molecules can activate Nrf2 signaling pathway leading to upregulation of cytoprotective genes such as detoxification and GSH biosynthetic enzymes. Under physiological condition, Nrf2 locates in cytoplasm, forming a protein complex with its cytosolic repressor Keap1.\textsuperscript{20,22} When cells are stimulated by oxidative stress and/or electrophiles, Nrf2 is disassociated from Keap1, followed by nuclear translocation. In the nucleus, Nrf2 forms a protein complex dimer with small Maf protein and other cofactors to control the expression of its target genes. In other words, translocation of Nrf2 can enhance the transcription of numerous antioxidative and Phase II detoxifying genes such as HO-1, NQO1, and GSTs.\textsuperscript{20,22} Thus, activation of Nrf2 signal can protect cells from challenges of drugs, toxicants, and carcinogens. Our results showed that HO-1, NQO1, and GCLM were upregulated by SCE at a transcriptional level in a concentration-dependent manner. In addition, SCE remarkably increased the total content of GSTs. Moreover, the SCE-mediated upregulation of HO-1 and NQO1 was abolished by Nrf2 knockdown. These results suggest that SCE modulated DMEs and drug transporters via the activation of the Nrf2 signaling pathway (Figure 12).

**Figure 12** Proposed underlying mechanism for the effect of SCE on drug metabolizing enzymes and drug transporters via Nrf2-mediated signaling pathway. Taken together, our findings have shown that pretreatment of human hepatocellular liver carcinoma cell line cells with SCE increased Nrf2 stability and half-life, mainly via downregulation of Keap1 and inhibition of Nrf2 degradation.

**Abbreviations:** SCE, Schisandra chinensis extract; Nrf2, nuclear factor (erythroid-derived 2)-like 2; Keap1, Kelch-like ECH-associated protein 1; ARE, antioxidant response element; NQO1, NAD(P)H:quinone oxidoreductase 1; HO-1, heme oxygenase-1; GSTA4, glutathione S-transferase A4; GCLM, glutamate–cysteine ligase, modifier subunit; GSTs, glutathione S-transferases; P-gp, P-glycoprotein; MRPs, multidrug resistance-associated proteins; OATP, organic anion transporting polypeptide.
Disassociation of Nrf2–Keap1 complex and translocation of Nrf2 into the nucleus for ARE binding in the promoter regions of target genes are the prerequisites of transcriptional activation of DME and transporter genes. In our experiments, we found that SCE concentration dependently enhanced Nrf2 translocation as well as increased total Nrf2 expression level. Our results also showed that SCE significantly inhibited Keap1 expression in a concentration-dependent manner. Previous studies have shown that Nrf2 activators might stabilize Nrf2 by increasing the ubiquitination and proteasomal degradation of Keap1 and consequently increased the half-life of Nrf2. In agreement with these findings, our findings showed that pretreatment of HepG2 cells with SCE increased Nrf2 stability and half-life mainly via downregulation of Keap1 and inhibition of Nrf2 degradation.

In this study, we found that both SchA and SchB induced the gene expression of HO-1 and NQO1 and prolonged the half-life of Nrf2 in HepG2 cells. However, SchA and SchB showed different potency and regulatory effects of some selected genes. In the SCE we prepared, its total lignanoid concentration was 44.97 mg/g crude extracts. In 300 μg/mL SCE used for cell treatment, the total lignanoid concentration is only 13.15 μg/mL, and thus the concentration of SchA and SchB in SCE was much lower than that we used for SchA at 20 μM and SchB at 200 μM. Our results demonstrate that the major components in SCE are lignanoids, and interaction of these lignanoid components may enhance the biological activities of SCE.

SCE at 100–300 μg/mL considerably up- or downregulated the expression of DMEs and drug transporters in HepG2 cells, but there is no clear dose response relationship, while maximum effect was often observed at 200 μg/mL SCE. The reason for this is unknown, but we suspect that gene regulation by SCE in HepG2 cells may be saturated when SCE concentration achieved 200 μg/mL, and the presence of multiple active components in SCE may antagonize and synergize depending on individual component concentrations. It has been reported that a low concentration of SCE exerts an antioxidative effect, whereas a high concentration of SCE exhibits an opposite effect. In the present study, we observed that SCE, SchA, and SchB induced the expression of NQO1, HO-1, and MRP2 in HepG2 cells, but this effect could not be fully abolished using Nrf2 siRNA. This suggests that there may be other related signaling pathways involved in the regulatory effect of SCE on the expression of DMEs and drug transporters in HepG2 cells. Recently, it has been reported that inhibition of c-Jun N-terminal kinase signaling pathway is involved in the liver protective effect of SCE in mice with acetaminophen-induced liver injury. Moreover, SCE or its components can inhibit nuclear transcription factor-κB and mitogen-activated protein kinase signaling pathways, contributing to its beneficial effects. Moreover, the RNA interfering efficiency may change, contributing to these varying effects.

**Conclusion**

In summary, our study has shown that SCE upregulates the expression of NQO1, HO-1, GCL, GSTA4, P-gp, and MRPs but downregulates the expression of OATPs primarily via the activation of Nrf2-mediated signaling pathway in HepG2 cells. Our findings may provide better understanding of the molecular mechanism for the hepatoprotective action of SCE in clinical practice. On the other hand, SCE may have potential to interact with other drugs, and this should be taken into consideration in polypharmacy. More mechanistic studies are needed to dissect how SCE protects the liver through regulation of cytoprotective DMEs and drug transporters.

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**Author contributions**

YY and SFZ designed and coordinated the study. JLH, ZWZ, JJY, and CQH conducted the experiments and data analysis. JLH and ZWZ drafted the manuscript. All authors critically reviewed the manuscript and approved the final version for publication.

**Disclosure**

The authors have no financial interests that might be perceived to affect the results or the discussion reported in this article.

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