The anticancer flavonoid chrysin induces the unfolded protein response in hepatoma cells

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Abstract

Chrysin is a natural and biologically active flavonoid with anticancer effects. However, little is known about the adaptive response of cancer cells to chrysin. Chrysin reportedly has proteasome inhibitor activity. Previous studies demonstrated that proteasome inhibitors might induce endoplasmic reticulum (ER) stress response. In this study, we aimed to determine the effects of chrysin on hepatoma cells and roles of the ER-resident protein GRP78 (glucose-regulated protein 78) in its action. Also, we investigated the effects of green tea polyphenol (-)-epigallocatechin-3-gallate (EGCG), a natural GRP78 inhibitor, on the sensitivity of hepatoma cells to chrysin. Here, we report that chrysin inhibits hepatoma cells growth and induces apoptosis in a dose-dependent manner. Chrysin induces GRP78 overexpression, X-box binding protein-1 splicing and eukaryotic initiation factor 2 \(^/H\) phosphorylation, hallmarks of the unfolded protein response. GRP78 knockdown potentiates chrysin-induced caspase-7 cleavage in hepatoma cells and enhances chrysin-induced apoptosis. EGCG overcomes chrysin-induced GRP78 expression. Combination of EGCG potentiates chrysin-induced caspase-7 and poly (ADP-ribose) polymerase (PARP) cleavage. Finally, EGCG sensitizes hepatoma cells to chrysin through caspase-mediated apoptosis. These data suggest that chrysin triggers the unfolded protein response. Abrogation of GRP78 induction may improve the anticancer effects of chrysin. Combination of EGCG and chrysin represents a new regimen for cancer chemoprevention and therapeutics.

Keywords: chrysin • endoplasmic reticulum stress • EGCG • glucose-regulated protein 78 • hepatoma

The stress-inducible glucose-regulated protein 78 (GRP78) is a multifunctional protein overexpressed in a variety of cancer cells [1]. In progressively growing tumours, GRP78 is highly induced by endoplasmic reticulum (ER) stress stimuli such as hypoxia, glucose starvation and low pH. Overexpression of GRP78 can stimulate tumour cells proliferation, survival, invasion and metastasis [2]. As an ER-resident chaperone, GRP78 interacts with transmembrane ER stress sensors such as inositol restriction enzyme 1, pancreatic ER kinase (PERK) and activation transcription factor 6 and control their activation; maintain Ca\(^{2+}\)/H\(^{11001}\) homeostasis and target misfolded proteins for proteasomal degradation [3]. Moreover, GRP78 may help to maintain ER integrity and inhibit ER-stress-induced apoptosis by preventing the activation of several pro-apoptosis molecules such as caspase-4, caspase-7 and Bik [4–6]. ER stress-induced autophagy is also dependent on GRP78 [7]. In addition to responding to the pathophysiological insults that disturb ER homeostasis, GRP78 confers resistance to chemotherapeutic drugs such as adriamycin, etoposide, temozolomide and fluorouracil [5, 8]. GRP78 expression can be induced by anticancer agents such as microtubules-targeting agents [9], Hsp70 inhibitor [10], histone deacetylase inhibitor [11] and anti-angiogenesis inhibitors [12, 13]. Blockade of GRP78 may sensitize cancer cells to these agents.

GRP78 is one of the targets for (-)-Epigallocatechin gallate (EGCG), a green tea polyphenol with cancer chemopreventive potential [14]. Mounting studies showed that EGCG inhibited the survival rate of malignant cells and induced apoptosis of malignant cells via the mitochondrial signal transduction pathway [15–17]. As a natural compound, EGCG is a promising agent for cancer chemoprevention. Also, EGCG may be of utility in cancer chemotherapeutics. Preclinical studies demonstrated that EGCG could sensitize tumour cells to temozolomide [8], quercetin [10], TNF-related apoptosis-inducing ligand (TRAIL) [18], paclitaxel and vincristine [9, 19]. EGCG is a natural inhibitor of GRP78 ATPase activity [14].

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As a GRP78 inhibitor, EGCG reportedly overcame resistance to ER stress-induced cell death in vitro [14]. In addition to GRP78, EGCG can target other molecules such as IGF-IR [20], Hsp90 [21] and BCL-2 [22], which play important roles in cell growth and survival. Chrysin (5,7-dihydroxyflavone) is a natural and biologically active flavonoid in many plants, honey and propolis. Previous studies have demonstrated that chrysin possesses potent anti-inflammatory and anti-oxidant properties [23, 24]. As an anti-oxidant, chrysin possesses vitamin-like activity in the body. Inhibition of the Cox-2 pathway is a mechanism underlying chrysin anti-inflammatory activity [24]. In light of these beneficial activities, chrysin has been described as disease-preventing dietary supplement. More importantly, chrysin may promote cancer cell death or perturb cell cycle progression [25, 26]. Although chrysin have activity as cancer-preventive agents, the molecular mechanisms underlying chrysin anticancer effects remain to be understood. Meanwhile, the elucidation of cellular adaptive response to chrysin may provide a novel strategy to sensitize cancer cells to this bioflavonoid. The combination of chrysin and other natural products or chemotherapeutic agents may be more effective in treating selective types of cancer.

Here, we provide evidence that chrysin inhibits hepatoma cell growth. The unfolded protein response is induced by chrysin in hepatoma cells. Chrysin induces the expression of GRP78. GRP78 knockdown sensitizes hepatoma cells to chrysin by potentiating chrysin-induced apoptosis. EGCG overcomes chrysin-induced GRP78 expression and enhances chrysin-induced cell death.

Materials and methods

Reagents

Chrysin and EGCG were purchased from MUST Biotech. (Chengdu, China). The caspase inhibitor z-VD-fmk was from Beyotime Institute of Biotechnology (Jiangsu, China). Anti-GRP78, anti-β-actin and anti-XBP-1 (X-box binding protein-1) antibodies were purchased from Santa Cruz Biotechnology (Santa Cruz, CA, USA). The anti-phosphorylated eukaryotic initiation factor 2α (eIF2α), anti-caspase-7 and anti-PARP antibodies were provided by Cell Signaling Technology, Inc. (Beverly, MA, USA).

Cell culture

Hepatoma cells HepG2 and SMMC-7721 were purchased from Cell Lines Bank, Chinese Academy of Science (Shanghai, China). Cells were grown in tissue culture flasks at 37°C in a humidified atmosphere of 5% CO2 and were maintained as monolayer cultures in DMEM supplemented with 5% foetal bovine serum and 100 U/ml penicillin and 100 μg/ml streptomycin.

Transfection of siRNA

The target sequence used for knockdown of GRP78 was 5’-GGAGCGCAU-UGAUACUAGA-3’. The negative control siRNA was purchased from Ribobio Co., Ltd. (Guangzhou, China). The double-stranded siRNA duplex was dissolved in DEPC-treated water. For transfection, 5000 cells were plated into 24-well plates and incubated for two days. LipofectAMINE 2000 reagent (Invitrogen, Carlsbad, CA, USA) was diluted in 50 μl of Opti-MEM I reduced serum medium and incubated at room temperature for 5 min. In addition, siRNA duplex was diluted in 50 μl of Opti-MEM I reduced serum medium and mixed with the pre-diluted LipofectAMINE 2000. The mixture was incubated at room temperature for 20 min., 50 nmol/l of siRNA was added into each well and incubated at 37°C.

Western blotting

Cells were washed twice with phosphate buffered saline and harvested with cold lysis buffer containing protease inhibitors. Cell lysates were collected from culture plates using a rubber policeman, and protein collected by centrifugation. Protein concentrations were determined by bicinchoninic acid (BCA) protein assay (Pierce Biotechnology, Rockford, IL, USA). Forty micrograms of total protein were boiled in 2× loading buffer (0.1 M Tris-Cl, pH 6.8, 4% SDS, 0.2% bromophenyl blue, 20% glycerol) for 10 min., then loaded into Tris-HCl-Polyacrylamide gels and transferred electrophoretically to Immobilon-P membrane (Millipore Corporation, Billerica, MA, USA). Membranes were incubated with primary antibodies and appropriate horseradish peroxidase (HRP)-linked secondary antibodies. Membranes were additionally probed with an antibody against actin to normalize loading of protein among samples. The secondary antibodies were detected by chemiluminescent agents (Pierce Biotechnology).

Cell viability assay

Hepatoma cells were plated in 96-well plates at 5000 cells per well. The next day, cells were treated with or without chrysin and EGCG in four replicates. After 48 hrs, cell viability was assessed by incubating cells with CCK-8 (Cell Counting Kit-8) reagents (Dojindo Laboratory Co., Ltd., Kumamoto, Japan) for 2–4 hrs and measuring the absorbance at 490 nm, and at 630 nm as reference, with a microplate reader (Bio-Rad, Hercules, CA, USA).

Cell apoptosis assay

Cell apoptosis was assessed by the Hoechst 33342 (Sigma-Aldrich, Inc., St. Louis, MO, USA) staining. Briefly, replicate cultures of hepatoma cells were plated in tissue culture plates. The cells were treated with or without chrysin and EGCG. After a change of fresh medium 24 or 48 hrs later, the cells were incubated with Hoechst 33342 solution at 37°C for 10 min., followed by examination under a fluorescence microscope. Strong fluorescence and condensed or fragmented nuclei can be observed in the nuclei of apoptotic cells, while weak fluorescence was observed in live cells. Quantification of apoptotic cells was performed by taking the images in random fields and counting at least 200 cells in four random fields in each well.

Statistical analysis

One-way ANOVA with least significant difference post hoc test was used to test for the differences in cell viability and apoptosis rate. All statistical tests were two-tailed, and difference to be considered statistically significant when P < 0.05.
Results

Chrysin inhibits hepatoma cell growth

To determine the effects of chrysin on hepatoma cell growth, HepG2 cells were treated with increasing doses of chrysin ranging from 2.5 to 40 μM for 48 hrs, followed by CCK-8 assay. Obvious change in cell shape was also detected after chrysin treatment. Whereas the untreated HepG2 cells displayed the cubic cell shape, many spindle cells were observed in chrysin-treated cells (Fig. 1A). Chrysin inhibited HepG2 cell growth in a dose-dependent manner (Fig. 1B). In addition, the inhibitory effects of chrysin on SMMC-7721 cells were observed, while SMMC-7721 cells were less sensitive to chrysin than HepG2 cells (Fig. 1C).

Next, we investigated whether chrysin induced hepatoma cell apoptosis. HepG2 cells were treated with increasing concentration of chrysin ranging from 2.5 to 40 μM for 48 hrs, followed by Hoechst 33342 staining. Although HepG2 cells growth was significantly inhibited by 2.5 μM chrysin (Fig. 1B), the apoptosis was not induced until the dosage of chrysin reached 10 μM. Increasing apoptosis rate was detected when the cells were treated with...
Chrysin at higher dosages (Fig. 2). These data suggested that chrysin inhibited cellular proliferation at relatively lower concentration, and induced apoptosis at relatively higher concentration.

**Chrysin induces the unfolded protein response**

Previous study indicated that chrysin possessed proteasome inhibitor activity [27]. Because proteasome inhibitor may induce ER stress, we investigated whether chrysin would induce ER stress or the unfolded protein response in cancer cells. HepG2 cells were treated with different doses of chrysin for 24 hrs, and then subjected to Western blot analysis. The results revealed that GRP78 expression was stimulated by chrysin in a dose-dependent manner (Fig. 3A). Similar effects were observed in another hepatoma cell line SMMC-7721 (Fig. 3A).

Next, we investigated the effects of chrysin on other ER stress responsive proteins including phosphorylated eIF2α and spliced XBP-1. During the unfolded protein response, a transient translation arrest is induced upon phosphorylation of eIF2α by PERK. The phosphorylation of eIF2α was observed after treatment of HepG2 cells with chrysin for 4 hrs (Fig. 3B). In addition, treatment of HepG2 cells with chrysin induced XBP-1 splicing, a hallmark of the unfolded protein response (Fig. 3C). Thus, chrysin is identified as an inducer of the unfolded protein response.

Caspase-7 is one of executioner caspases that mediate ER stress-induced apoptosis. To determine whether chrysin induce caspase-7 cleavage, HepG2 and SMMC-7721 cells were treated with different doses of chrysin for 24 hrs, followed by Western blot analysis of caspase-7 cleavage. Chrysin induced caspase-7 cleavage in a dose-dependent manner (Fig. 3D). In addition, treatment with chrysin induced PARP cleavage (Fig. 3E), a hallmark of cell apoptosis.

**GRP78 knockdown potentiates chrysin-induced apoptosis**

Because GRP78 represents a pro-survival arm in the UPR, down-regulation of GRP78 may break the balance of pro-survival signals and pro-apoptosis molecules. GRP78 can interact with caspase-7...
and prevent its activation. To investigate whether GRP78 down-regulation potentiates the activation of caspase-7 by chrysin, the effects of GRP78 knockdown by small interfering RNA on the cleavage of caspase-7 were examined. Treatment of HepG2 and SMMC-7721 cells with chrysin up-regulated GRP78 expression, but weakly induced caspase-7 cleavage. In contrast, GRP78 knockdown resulted in dramatic increase in chrysin-induced caspase-7 cleavage (Fig. 4). These data indicate that GRP78 blockade can potentiate the activation of caspase-7 by chrysin.

Next, we investigated the effects of GRP78 knockdown on chrysin-induced apoptosis. siRNA-mediated knockdown of GRP78 significantly enhanced chrysin-induced apoptosis in both HepG2 and SMMC-7721 cells. Treatment with caspase inhibitor abrogated the effects of chrysin and GRP78 knockdown on apoptosis (Fig. 5). These data suggested that GRP78 knockdown could sensitize hepatoma cells to chrysin by promoting caspase activation.

**EGCG overcomes chrysin-induced GRP78 expression and potentiates the activation of caspase-7 by chrysin**

Previous studies have identified EGCG, the major component of green tea, as a natural compound that overcomes paclitaxel-induced GRP78 expression and directly interacts with the ATP-binding domain of GRP78, blocks the interaction between GRP78 and procaspase-7, and suppresses the protective function of GRP78 [14, 19]. To investigate whether EGCG can overcome chrysin-induced GRP78 expression and enhance chrysin-induced caspase-7 activation, HepG2 cells were treated with EGCG, chrysin or both. Treatment with EGCG alone had no effects on GRP78 expression and caspase-7 cleavage. However, combination of EGCG suppressed chrysin-induced GRP78 expression and resulted in an increase in caspase-7 cleavage compared to that in cells treated with chrysin alone (Fig. 6A). Similar effects were observed in SMMC-7721 cells (Fig. 6A). EGCG also potentiated chrysin-induced PARP cleavage (Fig. 6B). These results demonstrated that EGCG could overcome chrysin-induced GRP78 expression and potentiate the pro-apoptotic signals induced by chrysin.

**EGCG sensitizes hepatoma cells to chrysin**

To determine whether combination of EGCG and chrysin could synergistically inhibit hepatoma cell growth, HepG2 and SMMC-7721 cells were treated with EGCG, chrysin or both for 48 hrs, followed by CCK-8 assay. Treatment of HepG2 and SMMC-7721 cells with 20 μM EGCG had little effects on cell growth. Treatment with 5 μM chrysin inhibited HepG2 cell growth. Combination of EGCG
Fig. 4 Potentiation of chrysin-induced caspase-7 cleavage by GRP78 knockdown. (A) HepG2 cells were transfected with negative control siRNA (siCtrl) or siRNA to GRP78 (siGRP78). Forty-eight hours later, the cells were treated with 10 μM chrysin for further 24 hrs. Total proteins were harvested and subjected to Western blot analysis of cleaved caspase-7, GRP78 and β-actin. (B) SMMC-7721 cells were transfected with negative control siRNA (siCtrl) or siRNA to GRP78 (siGRP78). Forty-eight hours later, the cells were treated with 20 μM chrysin for further 24 hrs. Total proteins were harvested and subjected to Western blot analysis of cleaved caspase-7, GRP78 and β-actin.

Fig. 5 Potentiation of chrysin-induced apoptosis by GRP78 knockdown. (A) HepG2 cells were seeded in a 24-well plate at 5000 cells per well. Two days later, cells were transfected with siRNA to GRP78 (siGRP78) or negative control siRNA (siCtrl). Forty-eight hours after transfection, the cells were treated with 10 μM chrysin for 48 hrs. Apoptosis was assessed by Hoechst 33342 staining. Red arrow: representatives of apoptotic cells. The apoptosis rate was plotted. Columns: mean percentage of apoptotic cells; bars: S.E. (B) SMMC-7721 cells were seeded in a six-well plate at 5000 cells per well. The next day, cells were transfected with siRNA to GRP78 (siGRP78) or negative control siRNA (siCtrl). Forty-eight hours after transfection, the cells were treated with 20 μM chrysin for 48 hrs. Apoptosis was assessed by Hoechst 33342 staining. Red arrow, representatives of apoptotic cells. The apoptosis rate was plotted. Columns: mean percentage of apoptotic cells; bars: S.E. A representative of two independent experiments was shown.
and chrysin yielded significantly more profound inhibition of cell growth than single agents (Fig. 7A). Whereas treatment of SMMC-7721 cells with 10 μM chrysin poorly inhibited cell growth, combination of EGCG and chrysin resulted in significant inhibition of SMMC-7721 cell growth (Fig. 7B).

To determine whether EGCG potentiated chrysin-induced apoptosis, HepG2 cells were treated with EGCG, chrysin or both for 48 hrs, followed by Hoechst 33342 staining. Whereas EGCG alone had little effects on apoptosis, combination of EGCG and chrysin resulted in significantly higher apoptotic rate than chrysin alone. The stimulatory effects of EGCG on chrysin-induced apoptosis could be abrogated by caspase inhibitor (Fig. 8A). EGCG also potentiated chrysin-induced apoptosis in SMMC-7721 cells (Fig. 8B). These data indicated that EGCG could potentiate chrysin-induced apoptosis through caspase activation.

Discussion

Previous studies have suggested that both flavonoids and green tea polyphenol can reduce the incidence of many types of cancer. Chrysin, a common flavonoid in plants, has been proposed as an antitumour agent. EGCG, the main constituent of green tea, also shows chemopreventive effects in animal models, as well as in epidemiological studies [27–29]. In this study, we have shown that chrysin inhibits hepatoma cell growth. Chrysin can induce ER stress response in hepatoma cells, including up-regulation of GRP78 expression, induction of eIF-2α phosphorylation and XBP-1 splicing. GRP78 knockdown potentiates chrysin-induced apoptosis. EGCG overcomes chrysin-induced GRP78 expression and synergistically promoted chrysin-induced apoptosis. These evidence may help to understand the mechanisms underlying the anticancer effects of chrysin.

Induction of ER stress is a mechanism underlying the anticancer effects of chemotherapeutic agents such as proteasome inhibitors and non-steroidal anti-inflammatory drugs [31, 32]. Upon ER stress, multiple signalling cascades may be activated to respond to perturbations in ER homeostasis. On one hand, ER stress response can promote cell survival through attenuation of protein synthesis and up-regulation of chaperones and folding enzymes [33]; on the other hand, prolonged or severe ER stress may ultimately overcome or bypass the cellular protective mechanisms, triggering cell death. We found that chrysin induced
GRP78 expression, eIF-2\alpha phosphorylation and XBP-1 splicing, hallmarks of the unfolded protein response. Chrysin is a known proteasome inhibitor [27]. Given that proteasome inhibition induces the unfolded protein response [31], it is reasonable to speculate that inhibition of proteasome may contribute, at least in part, to the induction of the unfolded protein response by chrysin. Tumour cells are generally more resistant to ER stress-induced apoptosis compared with un-transformed cells [34]. Overexpression of molecular chaperones in cancer cells may confer resistance to ER stress [35]. For example, GRP78 is a multifunctional chaperone that can protect cells from ER stress-induced apoptosis [3]. Mechanistically, GRP78 interacts with caspase-4, caspase-7 and BIK, and inhibits their activation [4–6]. Chrysin can induce the activation of pro-apoptotic signals, such as caspase-7 and PARP cleavage. In agreement with the inhibitory effect of GRP78 on caspase-7 activation [5], GRP78 knockdown enhanced chrysin-induced activation of caspase-7. Furthermore, GRP78 knockdown potentiated chrysin-induced apoptosis. Therefore, it can be concluded that GRP78 may protect hepatoma cells from chrysin-induced apoptosis.

Up-regulation of GRP78 by chrysin may represent an adaptive response. However, in higher concentrations like 40 to 60 \mu M of chrysin, caspase-7 cleavage and apoptosis was still observed in the presence of abundant GRP78 induction, suggesting that the
protective effects of GRP78 may be overcome or compromised by
deterious effects in the higher dosages. This is in agreement with
the scenario that the protective branches in ER stress responses
may attenuate but not completely abolish cell death upon severe
or prolonged ER stress. Severe ER stress may ultimately over-
come or bypass the cellular protective mechanisms, triggering cell
death. Nonetheless, blockade of the protective arms of ER stress
response may maximize the anticancer effects of chrysin. This is
in consistent with previous studies that show GRP78 blockade can
sensitize tumour cells to etoposide, temozolomide, quercetin and
paclitaxel [8–10, 14].

The combination of natural products in cancer chemoprevention
and therapy may prove to be more effective than single
agents. Given that cancer cells is capable of eliciting robust
response to resist exogenous insults including chemotherapeutic
agents, down-regulation of the protective stress proteins may
improve the therapeutic efficacy. The combination of a compound
that may induce stress proteins and another compound that
antagonizes these stress proteins may have synergistic effects on
cancer therapy. One example of this scenario is Hsp90 inhibitor
17-Allylamino-demethoxy geldanamycin (17-AAG). 17-AAG
inhibits the association of hsp90 with the heat shock factor-1
(HSF-1) and thereby rescues the inhibition of HSF-1 by Hsp90,
which leads to Hsp70 overexpression [36, 37]. Hsp70 may inhibit
several steps of the apoptotic cascades including release of
cytochrome C and apoptosis-inducing factor from the mitochon-
dria, nuclear import of AIF, activation of procaspases-9 and -3 and
even downstream of active caspase-3 [38–41]. Abrogation of
Hsp70 induction significantly enhances the anti-leukaemia activity
of 17-AAG [42]. The green tea polyphenol EGCG is known as an
inhibitor of stress protein GRP78 [14]. Combination of EGCG with
chemotherapeutic agents that induce the unfolded protein
response can enhance tumour cells sensitivity in these agents [8,
9, 14]. EGCG also synergistically promotes quercetin-induced cell
death [10]. In this study, we found that EGCG overcame chrysin-
induced GRP78 expression. These results suggest that EGCG may
have dual effects on GRP78, including inhibition of the ATPase
activity of GRP78 and inhibition of stress-induced GRP78 expres-
sion. Hence, combination of EGCG and chrysin may be more
effective than single agents. Indeed, EGCG sensitizes hepatoma
cells to chrysin-induced apoptosis, which is associated with
increased caspase and PARP cleavage. Thus, we speculate that
combination of chrysin and EGCG may be a novel regimen for can-
cer prevention or therapeutics.

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Conflict of interest

The authors confirm that there are no conflicts of interest.

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