Genistein reduces the activation of AKT and EGFR, and the production of IL6 in cholangiocarcinoma cells involving estrogen and estrogen receptors

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Abstract. Cholangiocarcinoma (CCA) is a malignant tumor of the biliary epithelium associated with Opisthorchis viverrini, primary sclerosing cholangitis and hepatitis viral infection. In the global population, men have higher incidence rates for CCA than women; thus, a gender disparity in the progression of chronic inflammation of the biliary duct leading to malignancy may involve the effects of estrogen (E2). Genistein (GE), a prominent phytoestrogen found in soy products, is an estrogen receptor β (ERβ) agonist and a tyrosine kinase inhibitor. The present study investigated the effects of GE on the growth of CCA cells by cell viability assay. The effects on signaling proteins were detected by western blot analysis and ELISA. Gene expression was examined by RT-qPCR. Two human intrahepatic CCA cell lines, HuCCA-1 and RMCCA-1, were utilized. GE (50-200 µM) reduced the viability of the two cell lines, and also inhibited the activation of epidermal growth factor receptor (EGFR) and AKT, as evidenced by decreasing protein levels of phosphorylated (p)-EGFR (Tyr1173) and p-AKT (Ser473), respectively. GE altered the mitogen-activated protein kinase signaling cascade by mediating decreased protein levels of p-p38 and increased protein levels of p-ERK1/2. GE significantly decreased the levels of interleukin 6 (IL6) and induced the expression of inducible nitric oxide synthase (iNOS). GE also downregulated the expression of p-ERα (Ser118) protein and ERα mRNA levels. Finally, GE induced the downregulation of the protein levels of ERβ. Of note, E2 deprivation potentiated the GE-induced reduction of p-EGFR (Tyr1173) and total AKT proteins and production of IL6, and mediated the downregulation of GE-induced iNOS protein. In conclusion, GE inhibited the growth of human CCA cell lines by reducing the activation of EGFR and AKT, and by attenuating the production of IL6. E2 and ER were also involved in the growth-inhibitory effect of GE in CCA cells.

Introduction

Cholangiocarcinoma (CCA) is one of the leading types of cancer in Thailand, with patients having a poor prognosis and treatment outcome. As there is no effective chemotherapeutic treatment at present, additional therapeutic approaches with specific and effective targets are required for CCA treatment. In Thailand, the incidence rate of CCA is associated with the prevalence of Opisthorchis viverrini infection (1). Chronic inflammation, often coupled with injury of the bile duct epithelium, is a common and important contributor to the malignant transformation of cholangiocytes. Nitric oxide, activated by inflammatory cytokines, is implicated in inflammation and the carcinogenesis of CCA (2). Interleukin 6 (IL6) is among the inflammatory cytokines that have been associated with the pathogenesis of CCA. The levels of IL6 have been observed to be elevated in patients with CCA, compared with those in healthy controls (3). Through the AKT signaling pathway, IL6 upregulates myeloid leukemia 1, resulting in resistance to apoptosis in CCA (4). The involvement of other signaling molecules has also been investigated in CCA cell growth and survival.

Several studies have reported the involvement of phosphatidylinositol-3 kinase (PI3K)/AKT/mammalian target of rapamycin (mTOR) signaling in CCA. It has been shown that the expression and activation of AKT protein increase during the development and progression of CCA (5,6), and that AKT is a promising target for CCA treatment (7). In addition, an activated PI3K/AKT signaling pathway is frequently found in CCA cells resistant to chemotherapy (8).

Epidemiological investigations have revealed gender disparities in CCA, for example, the incidence rate is higher in men (9). Liver biopsies from patients with intrahepatic CCA are positive for the estrogen receptor (ER), both ERα and ERβ subtypes, whereas normal liver cholangiocytes are ER-negative (10). The
growth-stimulating effects of estrogen (E2) via ER signaling or crosstalk with the epidermal growth factor receptor (EGFR) and AKT pathways are well established in breast cancer and endometrial cancer (11-14). E2 has also been shown to be involved in the proliferation (10,15) and invasion (16) of CCA. Additionally, E2 stimulates the production of IL6 in biliary epithelial cells (17). These observations suggest that the disruption of E2 may be useful for CCA therapy.

Genistein (GE), an isoflavone found in soybeans, has multi-targeted biological effects in cancer cells, leading to the inhibition of cell growth and induction of apoptotic cell death in several cancer cell lines, including breast (18), lung (19) and colon cancer (20). In addition to being a phytotoxin, the biological effects of GE have been reported to be associated with tyrosine kinase inhibition, anti-inflammatory effects (21,22) and the inhibition of AKT (23,24).

The present study investigated the effects of GE and the involvement of E2 in human CCA cell lines by targeting AKT, EGFR, ER, and IL6. The results showed that GE exhibited cytotoxicity in human HuCCA-1 and RMCCA-1 cell lines. GE reduced the protein levels of p-AKT and p-EGFR, and the production of IL6. Additionally, GE induced downregulation of the expression of ERα at the mRNA and protein levels. The present study demonstrated for the first time, to the best of our knowledge, that E2 deprivation potentiated the GE-induced reduction in the production of IL6, and protein levels of total AKT, p-AKT (Ser473) and p-EGFR (Tyr1173) in CCA.

Materials and methods

Chemicals and reagents. GE, 17-β estradiol (E2) and triciribine were purchased from Sigma-Aldrich; EMD Millipore (Billerica, MA, USA). LY294002 was obtained from Cayman Chemical Company (Ann Arbor, MI, USA). AG1478 was acquired from Merck Millipore (Darmstadt, Germany). Ham’s F12 medium, phenol red-free RPMI-1640 medium, and supplements were purchased from Gibco; Thermo Fisher Scientific, Inc. (Waltham, MA, USA). Fetal bovine serum (FBS) treated with dextran-charcoal (CSS) was purchased from HyClone; GE Healthcare Life Sciences (Logan, UT, USA). Antibodies were obtained as follows: Anti-human p-AKT (Ser473) cat. no. 13038, AKT cat. no. 9272, p-p38 (Thr180/Tyr182) cat. no. 9215, p38 cat. no. 9217, p-ERK1/2 (Thr202/Tyr204) cat. no. 9106 and ERK1/2 cat. no. 9102 from Cell Signaling Technology, Inc. (Danvers, MA, USA); ERα cat. no. sc-8002, p-ERα (Ser118) cat. no. sc-12915, p-EGFR (Tyr1173) cat. no. sc-12351 and EGFR cat. no. sc-03 from Santa Cruz Biotechnology, Inc. (Santa Cruz, CA, USA); ERβ cat. no. 05-824 and inducible nitric oxide synthase (iNOS) cat. no. ABN26 from Merck Millipore, and β-actin cat. no. A5316 from Sigma-Aldrich; EMD Millipore (Billerica, MA, USA). HRP-conjugated secondary antibodies obtained were as follows: Anti-rabbit IgG (cat. no. 5127) from Cell Signaling Technology, Inc.; anti-goat IgG (cat. no. HAF109) from R&D System Inc. (Minneapolis, MN, USA); and anti-mouse IgG (cat. no. 1706516) from Bio-Rad Laboratories, Inc. (Hercules, CA, USA).

Cell lines and cell culture. The HuCCA-1 and RMCCA-1 human intrahepatic CCA cell lines were established and kindly provided by Professor Stitaya Sirisinha and Dr Kawin Leelawat, respectively (25,26). The cells were cultured in complete medium (FBS condition) containing Ham’s F12 culture medium supplemented with 10% FBS, 2 mM L-glutamine, 100 U/ml penicillin and 100 µg/ml streptomycin. For the E2-deprived condition (CSS condition), cells were switched to a culture of phenol red-free RPMI-1640 medium supplemented with 10% CSS, 2 mM L-glutamine, 100 U/ml penicillin and 100 µg/ml streptomycin for 24 h prior to treatment. The cells were maintained at 37°C in a humidified atmosphere of 5% CO₂.

The difference between the FBS and CSS cultures was the concentration of steroid hormone E2. The E2 concentration in FBS is reported to be >100 pg/ml (27), whereas the concentration in CSS is in the range of 15-20 pg/ml (from COA of product). This model has been commonly used by others and by our group at the Laboratory of Pharmacology, Chulabhorn Research Institute (Bangkok, Thailand) as in vitro E2-deprived cell culture condition (28,29).

Cell viability assay. The cells maintained in the FBS condition were plated into 96-well plates at 1x10⁴ cells/well. Following incubation for 24 h, the medium containing various concentrations of GE (50-200 µM) or vehicle (0.1% DMSO) were added in triplicate wells and incubated at 37°C for 24 or 48 h. Following the manufacturer’s protocol, following this period of incubation, 10 µl of PrestoBlue™ cell viability reagent (Invitrogen; Thermo Fisher Scientific, Inc.) was used to detect cell viability. The reaction was measured by spectrofluorometry to detect fluorescence at 560 nm for excitation and 590 nm for emission using a SpectraMax microplate reader (Molecular Devices LLC, Sunnyvale, CA, USA).

Western blot analysis. The HuCCA-1 or RMCCA-1 cells were plated on a 100-mm culture dish at 4x10⁶ cells with 10 ml per plate and treated with GE (50, 100 and 200 µM for 0.5, 6 or 24 h), LY294002 (20 µM for 0.5, 6 or 24 h), triciribine (20 µM for 0.5, 6 or 24 h), or AG1478 (0.01 µM for 6 h) in either FBS or CSS conditions, according to each experimental design. Following the exposure period, the cells were collected and centrifuged 500 x g for 10 min at 4°C. The pellets were then re-suspended in RIPA buffer containing 50 mM Tris (pH 7.4), 150 mM NaCl, 1 mM EDTA, 0.1% SDS, 1% sodium deoxycholate, 1% NP40 and protease inhibitors (Calbiochem; EMD Millipore) mixed with 100 mM PMSF and phosphatase inhibitors (1 mM Na3VO4, 100 mM NaF and 1 nM okadaic acid). The protein concentration was determined using Bradford reagent according to the manufacturer’s protocol (Bio-Rad Laboratories, Inc.). For immunoblotting, 50 µg of total protein was separated using 10% SDS-PAGE and then electro-transferred onto a nitrocellulose membrane (Amersham Biosciences, Buckinghamshire, UK). The membrane was incubated in blocking buffer [5% non-fat dry milk in TBST buffer (10 mM Tris-HCl pH 8.0, 150 mM NaCl, and 0.05% Tween-20)] for 1 h at room temperature. The membrane was probed overnight with primary antibody p-AKT (Ser473) (1:1,000), AKT (1:2,000), p-p38 (Thr180/Tyr182) (1:2,000), p38 (1:2,000), p-ERK1/2 (Thr202/Tyr204) (1:2,000), ERK1/2 (1:2,000), ERα (1:1,000), p-ERα (Ser118) (1:1,000), p-EGFR (Tyr1173) (1:1,000), EGFR (1:1,000), ERβ (1:2,000) or iNOS (1:1,000) at 4°C, followed by incubation with appropriate HRP-conjugated secondary antibody (1:3,000)
at room temperature for 2 h. Subsequently, the blot was reacted with a chemiluminescent substrate (Amersham; GE Healthcare Life Sciences, Chalfont, UK), and exposed to X-ray film (GE Healthcare Life Sciences). The protein levels were quantified by densitometry using ImageQuant TL 7.0 software (GE Healthcare Life Sciences). The expression level of each protein was normalized to the corresponding protein level of β-actin derived from the same blot.

**Enzyme-link immunosorbent assay (ELISA).** The culture medium collected following treatment was analyzed for levels of IL6 using the DuoSet ELISA Development kit (R&D System Inc.). The absorbance was immediately measured at 450 nm, with the wavelength correction at 570 nm, on a SpectraMax microplate reader (Molecular Devices LLC., San Jose, CA, USA).

**Reverse transcription-quantitative polymerase chain reaction (RT-qPCR) analysis.** Total mRNA was isolated from 4x10^6 HuCCA-1 cells using an RNA purification cell kit (5 PRIME GmbH, Hamburg, Germany) according to the manufacturer's recommendations. Contaminating DNA was removed using the Turbo DNA-free™ DNase kit (Applied Biosystems; Thermo Fisher Scientific, Inc.). A master mixture was created in a 50 µl total reaction volume containing 1X RNA-direct™ SYBR®-Green Realtime PCR Master Mix (Toyobo Co., Ltd., Osaka, Japan), 0.2 µM forward primer, 0.2 µM reverse primer and 250 ng/µl RNA sample. The RT-qPCR analysis was performed on an ABI-StepOnePlus thermocycler (Applied Biosystems; Thermo Fisher Scientific, Inc.), starting with the RT reaction (10 min at 50˚C), followed by 40 cycles of two-step PCR (95˚C for 10 sec and 60˚C for 30 sec). Standard melting curve analysis was performed. The housekeeping gene, glyceraldehyde-3-phosphate dehydrogenase, was used for normalization. The relative quantification of the target genes was performed using StepOnePlus™ software version 2.1 (Applied Biosystems; Thermo Fisher Scientific, Inc.), based on the 2^ΔΔCq method, and resulted in the relative transcription level of the target genes in treated cells, compared with those in control cells. The primer sequences are listed in Table I.

**Statistical analysis.** Data are expressed as the mean ± standard error of the mean calculated from at least three independent experiments. Multiple comparisons were performed using one-way analysis of variance with Dunnett's multiple comparisons test by Stata™ program version 10.1 (StataCorp, College Station, TX, USA). P<0.05 was considered to indicate a statistically significant difference.

**Results**

GE reduces HuCCA-1 and RMCCA-1 cell viability. Two human CCA cell lines, HuCCA-1 and RMCCA-1, were used to investigate the cytotoxic effect of GE. As shown in Fig. 1,
GE concentrations ranging between 10 and 200 µM exhibited a dose- and time-dependent induction of cytotoxic effects in the HuCCA-1 and RMCCA-1 cells. A statistically significant cytotoxic effect (P<0.05) of GE was observed, starting at a GE concentration of 50 µM in the HuCCA-1 cells (Fig. 1A) and 100 µM in the RMCCA-1 cells (Fig. 1B) when treated for 48 h.

GE reduces the protein expression of p-AKT in HuCCA-1 and RMCCA-1 cells. AKT protein is a promising target for CCA treatment as its expression and activation increase during the development and progression of CCA (5-7). Studies in colorectal cancer cells have demonstrated that GE can induce apoptosis by inhibiting the activation of AKT (30). Therefore, the present study examined whether GE attenuated the activity of AKT in the CCA cell lines. LY294002 (PI3K inhibitor) and triciribine (AKT inhibitor) were used as positive controls. As shown in Fig. 2A-F, GE downregulated the protein expression of p-AKT (Ser473) in the two cell lines.

GE alters EGFR signaling in HuCCA-1 and RMCCA-1 cells. As AKT is one of the downstream molecules of EGFR, exhibiting prolonged activation in CCA (31), and GE is a known tyrosine kinase inhibitor (32), the present study aimed to determine the effect of GE on the protein expression of p-EGFR (Tyr1173) in the CCA cell lines. As shown in Fig. 3A-F, GE suppressed the protein expression levels of p-EGFR (Tyr1173) in the HuCCA-1 and RMCCA-1 cell lines.

The mitogen-activated protein kinases (MAPKs), both extracellular signal-regulated kinase (ERK)1/2 and p38, are involved in cell proliferation of KMCH-1 malignant cholangiocyte cells (33). Therefore, the present study investigated the effect of GE on ERK1/2 and p38. The results showed that GE decreased the protein expression levels of p-p38 (Thr180/Tyr182) but enhanced the protein levels of p-ERK1/2 in the two CCA cell lines, as shown in Fig. 4A-D.

Activation of AKT in HuCCA-1 cells is mediated by EGFR and other upstream molecules. The previous experiments...
demonstrated that GE induced a reduction of active p-EGFR and p-AKT. The activation of AKT can be mediated by upstream molecules other than EGFR. It has been shown that non-genomic ER signaling can crosstalk with EGFR, which leads to the activation of ERK1/2 and consequently induces cell growth (29). The present study examined whether the activation of AKT in CCA cells is mediated only by EGFR, or involves other molecules, including ER. The preliminary results revealed that HuCCA-1 cells expressed ERα and ERβ proteins, whereas RMCCA-1 cells expressed only ERβ protein. Therefore, only HuCCA-1 cells were utilized in this and the following experiments in order to investigate the role of ER in the effects of GE. To examine whether the GE-inhibited activation of AKT is mediated by EGFR or other molecules, the EGFR inhibitor (AG1478) was utilized as a positive control. As shown in Fig. 5A and B, GE markedly reduced the protein expression levels of p-EGFR (Tyr1173) and p-AKT (Ser473), whereas AG1478 (10 nM) inhibited the protein expression of p-EGFR (Tyr1173) only. These results suggested that, in addition to EGFR, other upstream molecules may be involved in the phosphorylation of AKT in HuCCA-1 cells.

Reduction of total AKT protein and inhibition of the activation of EGFR by GE are enhanced under conditions of E2 deprivation. The two CCA cell lines used in the present study are intrahepatic CCA cell lines. HuCCA-1 cells were originated from O. viverrini-infected patients, whereas RMCCA-1 cells were derived from a patient, whose parasitic infection information was not reported (25,26). Therefore, the two cell lines were used in the present study. E2 is involved in CCA cancer proliferation (10,15) and crosstalk exists between ER and EGFR signaling in breast cancer cell lines (29). Therefore, it was hypothesized that E2 may be partly involved in the activation of EGFR in CCA cells. In order to investigate the role of E2 in the effects of GE, experiments designed in CSS conditions were compared with those in FBS conditions. To confirm whether E2 is involved in the GE-induced activation of EGFR, HuCCA-1 cells were cultured and treated with GE under CSS conditions. As expected, GE markedly suppressed the protein expression level of p-EGFR (0.61±0.11-, 0.57±0.10- and 0.55±0.05-fold of control at GE concentrations of 50, 100, and 200 µM, respectively) under E2 deprivation (Fig. 6A). However, in FBS conditions (Fig. 3), GE reduced the protein expression level of p-EGFR (0.61±0.11-, 0.57±0.10- and 0.55±0.05-fold of control at GE concentrations of 50, 100, and 200 µM, respectively) under E2 deprivation (Fig. 6A).
Figure 4. Effect of GE on MAPK proteins in HuCCA-1 and RMCCA-1 cells. Western blot analysis of (A) p-ERK1/2 and ERK, and (B) p-p38 and p38 in HuCCA-1 cells treated with 50, 100 and 200 µM GE for 24 h in complete medium. Western blot analysis of (C) p-ERK1/2 and ERK, and (D) p-p38 and p38 in RMCCA-1 cells treated with 50, 100 and 200 µM GE for 24 h in complete medium. Quantification of the western blot data is shown in lower panels. Data are presented as the mean ± standard error of the mean of at least five independent experiments. *P<0.05, vs. C. GE, genistein; ERK, extracellular signal regulated kinase; C, control (0.1% DMSO vehicle); p-, phosphorylated.

Figure 5. Activation of AKT in HuCCA-1 cells is not only mediated by EGFR. Western blot analysis of (A) p-EGFR (Tyr1173) and EGFR, and (B) p-AKT (Ser473) and AKT, in HuCCA-1 cells treated with 0.1% DMSO vehicle control, 10 nM AG, and 50 µM GE for 6 h in complete medium (upper panels), and quantification of the western blot data (lower panels). Data are presented as the mean ± standard error of the mean of at least five independent experiments. *P<0.05, vs. C. GE, genistein; AG1478; C, control; EGFR, epidermal growth factor receptor; p-, phosphorylated.
expression levels of p-EGFR to a lesser degree (0.78±0.11-, 0.66±0.10- and 0.64±0.08-fold of control for GE concentrations of 50, 100, and 200 µM, respectively). Additionally, the down-regulation of total protein levels of AKT induced by GE under E2-deprivation (0.93±0.05-, 0.88±0.10- and 0.66±0.06-fold of controls for GE concentrations of 50, 100, and 200 µM, respectively; Fig. 6B) was higher, compared with the corresponding effect in FBS conditions (1.07±0.10-, 0.88±0.09- and 0.81±0.09-fold of control for GE concentrations of 50, 100, and 200 µM, respectively; Fig. 2). The inhibitory effects of GE on p-EGFR (Tyr1173) and total AKT were enhanced in the CSS condition, compared with those in the FBS condition. These findings supported the hypothesis that E2 and its receptors may be involved in the activation of EGFR, and may interfere with the reduced phosphorylation of EGFR induced by GE.

GE reduces the production of IL6 and alters the expression of iNOS in HuCCA-1 cells. The inflammatory cytokine IL6 is one of the therapeutic targets for CCA (34) and GE has been reported to inhibit the IL6 produced by macrophages (21). Previous experiments have revealed that GE reduces the activation of AKT, which is also a downstream molecule of the IL6 pathway. Therefore, the present study investigated the effect of GE treatment in HuCCA-1 cells on the production of IL6 and the inflammation-associated protein, iNOS. As shown in Fig. 7A, treatment with GE for 6 h decreased the level of IL6 in a dose-dependent manner in the FBS and CSS conditions. Of note, with the same concentration of GE, treatment in the CSS condition led to a more marked reduction in IL6, compared with that in the FBS condition. The fold changes from the controls in the FBS condition were 0.10±0.04, 0.25±0.02 and 0.41±0.07, whereas those in the CSS condition were 0.33±0.04, 0.54±0.04 and 0.66±0.05 at GE concentrations of 50, 100 and 200 µM, respectively. In the CSS condition, GE reduced the protein level of iNOS, whereas GE significantly increased the protein level of iNOS in the FBS condition (Fig. 7B).

Figure 6. Effect of GE on AKT and EGFR under E2 deprivation. Western blot analysis of (A) p-EGFR (Tyr1173) and EGFR, and (B) p-AKT (Ser473) and AKT, in HuCCA-1 cells treated with 50, 100 and 200 µM GE for 6 h in CSS conditions (upper panels), and quantification of the western blot data (lower panels). In CSS conditions, cells in culture plates were maintained in CSS condition 1 day prior to continual treatment with E2 or GE for 6 h. Data are presented as the mean ± standard error of the mean of at least five independent experiments. *P<0.05, vs. C. GE, genistein; E2, estrogen; CSS, E2-deprived conditions; C, 0.1% DMSO vehicle control in CSS; EGFR, epidermal growth factor receptor; p-, phosphorylated.

Figure 7. GE reduces induction of IL6 in HuCCA-1 cells. HuCCA-1 cells treated with 50, 100 and 200 µM GE for 6 h in FBS or CSS conditions. Following treatment, the cultured media were collected to determine the (A) protein level of IL6 by enzyme-link immunosorbent assay analysis. The level of IL6 (pg/ml) was normalized to the total protein of each sample (µg). The cell lysates were harvested to evaluate the expression level of (B) iNOS using western blot analysis (upper panel), with quantification of the western blot data (lower panel). Data are presented as the mean ± standard error of the mean of at least four independent experiments. *P<0.05, vs. C in FBS; #P<0.05, vs. C in CSS. GE, genistein; FBS, complete medium; CSS, E2-deprived conditions; C, 0.1% DMSO vehicle control; IL6, interleukin 6; iNOS, inducible nitric oxide synthase.
GE alters the protein and gene expression of ER in HuCCA-1 cells. The results of the present study showed that GE effectively downregulated the phosphorylation of EGFR and induction of IL6 in CSS, compared with FBS conditions, suggesting that ER may be involved in those effects. In addition, GE has been reported to reduce the expression of ER in rat prostates (35). Therefore, the present study further investigated the effects of GE treatment on the expression of ER in HuCCA-1 cells. As shown in Fig. 8A, GE dose-dependently downregulated the protein expression of ERα at 24 h in FBS and CSS conditions. In addition, the results demonstrated that treatment with GE (100 and 200 µM) for 24 h in the CSS condition significantly reduced the level of p-ERα (Ser118), whereas GE at a concentration of 50 µM increased the level of p-ERα (Ser118). Additionally, the GE-induced downregulation of ERβ was observed following 6 h of treatment in the FBS condition and following 24 h of treatment in CSS condition (Fig. 8B).

To obtain further insight into the effect of GE on ERα and ERβ in HuCCA-1 cells, RT-qPCR analysis of the mRNA expression of ERα (left) and of ERβ1 and ERβ2 (right) in HuCCA-1 cells treated with 50, 100 and 200 µM GE for 6 h in FBS or in CSS, with quantification of the western blot data (lower panels). Relative mRNA expression of ERα (left), and of ERβ1 and ERβ2 (right) in HuCCA-1 cells treated with or 50, 100 and 200 µM GE for 6 h in FBS or in CSS, analyzed using reverse transcription-quantitative polymerase chain reaction analysis. Data are presented as the mean ± standard error of the mean of at least three independent experiments. *P<0.05, vs. C in FBS; #P<0.05, vs. C in CSS. GE, genistein; FBS, complete medium; CSS, E2-deprived conditions; C, 0.1% DMSO vehicle control in FBS or CSS; ER, estrogen receptor; p-, phosphorylated.
The present study showed that GE significantly inhibited the activity of EGFR, as evidenced by a decrease in active p-EGFR (Tyr1173). Previous reports have shown that EGFR is involved in the proliferation and progression of CCA (31,41,42). It has been shown that GE suppresses the proliferation of proliferative cholangitis in rat models, where the expression of EGFR is high (43). Furthermore, GE conjugated with recombinant human EGF has the capability to inhibit EGFR tyrosine kinase and to trigger rapid apoptotic cell death in EGFR-positive MDA-MB231 and BT-20 breast cancer cells (44). These reports support the results of the present study that GE inhibited the activities of AKT and EGFR, and this was involved in the reduced viability observed in the HuCCA-1 and RMCCA-1 cells.

It is well established that MAPK signaling can be activated upon the activation of EGFR. ERK1/2 and p38 are members of the MAPK family of cell signaling proteins. The signaling cascades of ERK1/2 and p38 and their upstream components, including ras, Raf, MEK and EGFR are associated with the transformation and progression of several types of cancer (45). A study investigating the KMCH-1 malignant cholangiocyte cell line indicated that mitogens can stimulate cell growth via the activation of ERK1/2 and p38. However, only p38 was involved in the anchorage-independent growth of this cell line (33). The present study revealed that GE inhibited the activation of p38, as evidenced by the reduction in the protein expression level of p-p38 in HuCCA-1 and RMCCA-1 cells. This finding is in line with a previous report in human prostate cancer cell lines that GE inhibited the activation of p38, leading to the inhibition of cell invasion (46). Therefore, it is possible that GE may also inhibit anchorage-independent transformed cell growth through the inactivation of p38 in CCA. Despite the reduced activation of p38, the results of the present study demonstrated that GE increased the protein expression levels of p-ERK1/2 in the two CCA cell lines. The activation of ERK1/2 is known to be involved in cell proliferation. However, several studies have demonstrated enhancement in the protein levels of p-ERK1/2 in cells undergoing apoptosis. For example, cisplatin increased the protein levels of p-ERK1/2 in a dose- and time-dependent manner, which was involved in the induction of apoptosis in HeLa cells (47). The GE-induced induction of G2/M cell cycle arrest in MDA-MB231 human breast cancer cells was also mediated via the activation of ERK1/2 (48). These reports are in line with the results of the present study, which showed that GE induced the phosphorylation of ERK1/2 in CCA cells. Taken together, the present study suggested that ERK1/2 and p38 were involved in the growth inhibitory effect of GE in CCA cells, but through a different mechanism of action. GE affects the inhibition of cell invasion and induction of cell cycle arrest through MAPK signaling. However, the present study did not focus on the effects of GE on cell cycle machinery. The effects of GE on MAPK signaling in CCA require further investigation.

AKT is the intermediate molecule of various pathways, including EGFR. Upon inhibiting EGFR kinase activity with AG1478, the activity of AKT is reduced, as demonstrated by a reduction in the phosphorylation of AKT in colorectal adenocarcinoma (49), nasopharyngeal carcinoma (50) and breast cancer cells (51). In the present study, although GE treatment...
led to a reduction in the phosphorylation of EGFR and AKT. Treatment with AG1478, an EGFR kinase inhibitor, in the HuCCA-1 cells reduced the phosphorylation of EGFR only, and not its downstream molecule AKT. These results suggested that, at least in HuCCA-1 cells, activation of the phosphorylation of AKT is induced not only by EGFR. Human epidermal growth factor receptor 2, another member of the EGFR family, has been shown to be involved in HuCCA-1 cell proliferation and invasion through the AKT pathway (52). In addition to the EGFR family of proteins, the activation of AKT by IL6 has also been reported. It has been shown that the growth of CCA cell lines stimulated by IL6 occurred via an AKT-dependent mechanism (53). Of note, in the present study, the suppression of p-AKT by GE was more prominent following a short (0.5 h) duration of treatment, compared with a longer (6-24 h) duration of treatment. This may be due to cell compensation for the reduced activity of AKT, which is a crucial molecule in several pathways, particularly those involved in cell survival and chemoresistance in CCA (5-8). Additionally, a previous study suggested that E2 enhances the protein expression levels of p-AKT, p-ERK1/2 and ERα, leading to the stimulation of CCA cell proliferation (10).

Previous studies have indicated that there is crosstalk between ERα and EGFR in breast cancer, leading to activation of the downstream molecules of EGFR signaling pathways, including MAPK and AKT pathways (12,29,31,54,55). Additionally, it has been reported that serum levels of E2 in patients with CCA increase in concentrations corresponding to an optimal dose for stimulating the proliferation and invasion of CCA cell lines in vitro (16). Based on the above information, and as GE is also an agonist of ERβ, the present study investigated whether E2 and ER were involved in the inhibitory effect of GE observed in the CCA cell lines. The results showed that the GE-induced inhibition of the activities of AKT and EGFR was enhanced in the absence of E2. This suggested the involvement of E2 in the inhibitory effect of GE and the growth of HuCCA-1 cells. In the presence of E2, GE competes with E2 for ER binding and, as it has a higher affinity towards ERβ, causes a growth inhibitory effect, which is a predominant effect of the activation of ERβ (56). In E2 deprivation, GE binds to ERβ, causing increased growth inhibition. However, the growth inhibitory effect of GE may be independent of the ER-mediated pathway. As reported in other cell lines and evidenced in CCA cells in the present study, GE possesses a tyrosine kinase-inhibitory effect. The GE-induced inhibition of the activities of EGFR and AKT may lead to a reduction of CCA cell growth. Therefore, the inhibition of EGFR and AKT evidenced following GE treatment may be mediated via its tyrosine kinase inhibiting effects and/or via interfering with the crosstalk between non-genomic ER and EGFR signaling.

CCA is associated with chronic inflammation of the bile duct (57). Upon binding to its receptor, E2, has been shown to act as an anti-inflammatory and anticarcinogenic by inhibiting the production of IL6 in DEN-induced liver cancer in animal models (58). The cytokine IL6 is important in CCA cell growth. Its signaling involves a complex network of different pathways, including the AKT pathways (4,59) and MAPK (60) pathways. The data presented in the present study showed that GE reduced the production of IL6 in HuCCA-1 cells. This result is in line with previous reports that GE possesses anti-inflammatory properties (22) and can inhibit the production of IL6 by macrophages stimulated with lipopolysaccharide (21). Additionally, the present study found that the GE-induced decrease in the production of IL6 in CSS conditions was more marked, compared with that in FBS conditions. These results suggested that E2 was involved in the phosphorylation of AKT, and also in the induction of IL6 in HuCCA-1 cells. This finding is in agreement with a previous report that E2 stimulated the production of IL6 in non-malignant biliary epithelial cells and malignant biliary epithelial cells expressing ERα (17). Therefore, in addition to EGFR, the results suggested that E2, ER and IL6 may also be involved in the inhibitory effect of GE and the growth of CCA cells.

Although it has been reported that GE can attenuate the nitric oxide released via iNOS in activated macrophages (61), the results of the present study demonstrated that GE increased the protein expression levels of iNOS in HuCCA-1 cells, despite decreased levels of IL6. However, a report by Nakaya et al demonstrated that GE and E2 upregulated the protein expression and activity of iNOS through ER pathways in RAW246.7 macrophage cells (62). This finding is in accordance with the results of the present study, which showed that GE induced the expression of iNOS in the presence of E2. The discrepancy in these findings may be due to differences in cell type and inflammatory stages. The HuCCA-1 cell line was established from CCA tissue fragments surgically removed from a male patient from Thailand diagnosed with chronic infection due to liver fluke (25).

The results of the present study suggested the involvement of E2 on the growth-inhibiting effects of GE in CCA cells. This suggests that its receptor, ER, is likely to be involved. GE has been identified as an agonist of ERβ; however, this compound can bind ERα and ERβ, of which binding affinity to ERβ is 20-30-fold higher than that to ERα (63,64). The data in the present study showed that, at 24 h, GE (100 and 200 μM) in CSS conditions reduced the protein expression levels of p-ERα (Ser118), ERα and ERβ, whereas the mRNA expression of ERα was downregulated at 6 h. This result is consistent with a previous report that showed that GE downregulated the expression of ER in rat prostates (35). In the present study, although GE reduced the protein expression of ERβ, there were no significant changes in the mRNA expression levels of ERβ1 and ERβ2 in the CSS conditions. These results suggested that the GE-induced reduction in ERβ protein was not at the transcriptional level. However, the effects of GE on p-ERβ were not measured in the present study, as information on the function of p-ERβ remains limited. As the ER pathway has been shown to be associated with the AKT (12) and IL6 (17) signaling cascades, the downregulation of ER induced by GE observed in the present study may be involved in the GE-induced inhibition of IL6 and AKT in CCA.

In conclusion, the present study is the first, to the best of our knowledge, to demonstrate that GE can reduce the growth of CCA cell lines via inhibition of the activities of EGFR, AKT and IL6. The mechanism was, in part, associated with downregulation of the protein and mRNA expression levels of ER. The present study provided preliminary results for further
investigations to generate further translational value for GE in the treatment of CCA, for example, modifying GE structure to improve its activity, using GE to sensitize CCA to conventional treatments, and potentially using GE for chemoprevention as a therapeutic supplement in CCA.

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Availability of data and materials

The analyzed datasets generated during the study are available from the corresponding author on reasonable request.

Authors' contributions

PT performed the experiments, data analysis and drafted the manuscript. AT initiated and monitored the research, supervised data interpretation and the drafting and revision of manuscript. PW and JS performed data interpretation and also assisted in the drafting of the manuscript. All authors have read and approved the final manuscript.

Ethics approval and consent to participate

Not applicable.

Consent for publication

Not applicable.

Competing interests

The authors declare that they have no competing interests.

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