Inhibitory Effect of Genistein on PLC/PRF5 Hepatocellular Carcinoma Cell Line

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ABSTRACT

Background: Natural compounds including flavonoids like genistein (GE) are able to inhibit cell proliferation and induce apoptosis. GE is the main representative of these groups. GE inhibits carcinogenic tumors such as colon, stomach, lung, and pancreas tumors. The aim of the present study was to analyze the apoptotic effect of GE in the hepatocellular carcinoma (HCC) PLC/PRF5 cell line.

Methods: Cells were treated with various doses of GE (1, 5, 10, 25, 50, 75, and 100 µM/L) at different times (24, 48, and 72 h) and the MTT assay was commonly used. Furthermore, cells were treated with single dose of GE (25 µM) at different times and flow cytometry was performed.

Results: GE inhibited the growth of liver cancer cells significantly with a time- and dose-dependent manner. The percentage of living cells in GE treatment groups with a concentration of 25 µM at different times were 53, 48 and 47%, respectively \((P < 0.001)\). Result of flow cytometry demonstrated that GE at a 25 µM concentration induces apoptosis significantly in a time-dependent manner. The percentage of apoptotic cells at different times were 44, 56, and 60%, respectively \((P < 0.001)\).

Conclusions: GE can significantly inhibit the growth of HCC cells and plays a significant role in apoptosis of this cell line.

Keywords: Apoptosis, genistein, hepatocellular carcinoma, proliferation

INTRODUCTION

Hepatocellular carcinoma (HCC), malignant hepatoma, is primary cancer of hepatocytes, main cell type of liver.\(^1\) This disease account for 85–90% of liver cancer and its survival is 6–20 months.\(^2\) HCC is considered as the eighth most frequently diagnosed cancer worldwide, and the third most frequent cause of cancer death\(^3\) and annually causes...
about 600/000 death.\[^4\]\ It is clinically asymptomatic until advanced stage and has a poor prognosis, low survival and high rate of recurrence.\[^5\]\ The disease has not uniformly distribution.\[^6\]\ There are three geographical areas including low, moderate, and high recurrence areas.\[^7\]\ More than 80% of the cases have been reported in South Africa and West Asia.\[^8\]\ The incidence of the disease is moderate in France, Germany and Great Britain and low prevalence in North and South America and North Europe.\[^9\]\ There are large geographical correlation between the incidence of the disease and hepatitis B. Many factors such as age and sex can affect the prevalence of the disease\[^9\]\ and morbidity of the men is more than the woman.\[^10\]\ HCC is the sixth most common cancer in men and the eleventh most common cancer in women.\[^11\]\ Cirrhosis is the final stage of chronic diffuse liver disease\[^11\]\ which in 5% of patients lead to HCC.

The main causes of cirrhosis are alcohol, hepatitis B, and hepatitis C. Infection with HBV and HCV are the major risk factors of this disease.\[^11\]\ Other risk factors include alcohol, tobacco, obesity, diabetes mellitus, inherited and metabolic disease. Among all natural compounds, flavonoids with anticancer activity have pharmacological activities such as antioxidant, anti-mutagenic, anti-bacterial, anti-angiogenic and anti-inflammatory effects. These compounds are often found in cereals such as soybeans and soybean products. Genistein (GE), a hydroxyisoflavone with a heterocyclic and diphenolic structure similar to estrogen, is the main representative of these groups.\[^12\]\

Genistein inhibits carcinogenic tumors such as colon, stomach, lung, and pancreas tumors (Andres et al., 2011). Ovarian cancer, breast, and prostate cancer in Asian countries where soy foods are high is lower than in Western countries\[^13\]\ and also consumption of soya foods is associated with reduced risk of HCC. It is important to note that 30–40% of cancers are because of dietary choices\[^14,15\]\ which are preventable by consumption of suitable fruits, soybean, and vegetables appropriate diets.\[^15,16\]\ Natural compounds including flavonoids such as quercetin,\[^17\]\ resveratrol, epigallocatechin gallate, lycopene, ursolic acid, isothiocyanates, perillyl alcohol,\[^18\]\ and curcumin are able to inhibit cell proliferation and induce apoptosis. Other natural compounds with anticancer effect are apigenin, GE, and daidzein.\[^19\]\ Quercetin has an antiproliferative effect on colon carcinoma (HCT-116 and HT-29) and mammary adenocarcinoma (MCF-7) cell lines\[^20,21\]\ and also apigenin inhibits proliferation in MCF-7 cells and HL-60 cells.\[^22,23\]\ GE has a preventive effect on breast cancer. GE induces apoptosis and inhibits growth of human leukemic MOLT-4 and HL-60 cells.\[^24,25\]\ Numerous studies have demonstrated that intake of high flavonoids diet decreases colon cancer and breast cancer\[^26-28\]\ and GE induces apoptosis in human pro-myelocytic HL-60 leukemic cells, prostate cancer (PCa, LNCGP, DU-145, PC-3) cells, and H-460 nonsmall lung cancer cells in a dose-dependent manner.\[^29-33\]\ It has been indicated that soy isoflavones (GE, genistein, daidzein, and biochanin A) inhibit growth of murine (MB-49 and MBT-2) and human (HT-1376, UM-UC-3, IF-4, J-82 and TCCSUP) bladder cancer cell lines in a dose-dependent manner. Epidemiological studies have shown that prostate cancers in Asians is lower than Americans because of consumption of soy products.\[^32,33\]\ More than 100 studies have indicated that GE has an inhibitory effect on various cancers such as melanoma, lung, leukemia, lymphoma, and bladder cancer.\[^36\]\

Apoptosis, programmed cell death and vital component of various processes, is characterized by distinct morphological changes that occur during apoptosis (Hacker, 2000) including shrinkage and pyknosis (chromatin condensation). During the process of apoptosis, plasma membrane blebbing occurs followed by karyorrhexis and separation of cell fragments into apoptotic bodies during a process called “budding” which these bodies consist of cytoplasm with packed organelles with or without a nuclear fragment.\[^37\]\

However, GE compound are obtained from vegetables, fruits, spices, teas, herbs, and medicinal plants, such as flavonoids, carotenoids, phenolic compounds, and terpenoids which high intake of these natural compounds decrease risk of human malignancies, including colon, breast, lung, laryngeal, pancreatic, bladder, stomach, esophageal, and oral cancers.\[^38,39\]\

Finally, epidemiological and experimental studies demonstrated that GE has a significant role in the prevention of cancers like breast cancer, prostate cancer, colon cancer, leukemia, melanoma, etc.\[^40\]\ but only limited data are available to demonstrate the effects of GE on HCC cell lines such as Bel 7402 cells, herein, a study has reported that GE significantly inhibits the growth of Bel 7402 cells and induces apoptosis in this cell line but there isn’t any report about effect of GE on PLC/PRF5 cell line, therefore, we decided to investigate apoptotic effect of GE on PLC/PRF5 cell line.\[^41\]\

METHODS

Human HCC cells (PLC/PRF5) were purchased from the National Cell Bank of Iran-Pasteur Institute. GE, Dulbecco minimal essential medium (DMEM) and MITT (5-[1,2-thiazolyl]-2,5-diphenyl-3H-tetrazolium bromide) were purchased from Sigma (Sigma, St. Louis, MO, USA). All other chemicals were obtained from the best sources available.

Cell culture

The cells were cultured in DMEM with pH 7.2-7.4 (Sigma) containing 1% sodium pyruvate (sigma),...
3.7 mg/ml sodium bicarbonate (Sigma), 10% fetal bovine serum (sigma) and 1% antibiotics which include 10,000 units/ml penicillin G sodium (sigma), 10,000 µg/ml streptomycin sulfate and 25 µg/ml amphotericin B (sigma) at 37°C in 5% CO₂ to promote attachment. When cells became > 80% confluent, 5 × 10⁵ cells were seeded into 24-well plates (Becton-Dickinson) for 24 h in DMEM culture medium before they were incubated with certain concentrations of GE (1, 5, 10, 25, 50, 75, and 100 µM/L), which was dissolved in dimethyl sulfoxide (DMSO); DMSO was present at 0.01-0.3% in the medium based on IC50 index, at different times (24, 48, and 72 h). The control cells were treated with DMSO only. Photography was done for cultures before and after treatment with GE at different times using inverted microscope (Nikon, TE 2000-U, Japan).

Determination of IC50 value by MTT assay
The effect of GE on cellular proliferation was assessed by MTT assay according to standard protocols. After 24, 48, and 72 h of the treatment, the IC50 value for GE in PLC/PRF5 groups were determined. The MTT assay was commonly used to assess cell proliferation and viability by measuring the reduction of yellow MTT by mitochondrial dehydrogenases in viable cells. Briefly, 5 × 10⁴ Cells were counted and placed into each well of a 24-well microplate and were treated with various drug concentrations (1, 5, 10, 25, 50, 75, and 100 µM/L) of GE for 24, 48, and 72 h and the MTT survival assay was then carried out for the evaluation of the cell viability with different drug concentrations. The cells were measured spectrophotometrically at 570 nm. All experiments were repeated 3 times, with at least three measurements (triplicates).

Determination of cell viability by MTT assay
To determine the effect of GE, the cells were seeded in triplicate in 24-well plates and treated with GE at a concentration of 25 µM in different period times (24, 48, and 72 h). Cell viability was estimated by a colorimetric assay based on the conversion of tetrazolium dye (MTT) to a blue formazan product. The absorbance of the cell lysates in DMSO solution was read at 570 nm by a microplate reader (Bio-Rad Hercules, CA, USA).

Determination of apoptotic cells by flow cytometry assay
The cells were seeded in 24-well plates. After 24 h, the medium was changed, and medium contains GE (25 µM) was added. After 24:48 and 72 h of incubation, all the adherent cells were collected with 0.05% trypsin, washed with cold phosphate-buffered saline (PBS) and resuspended in binding buffer (×1). After addition of Annexin V-FITC and propidium iodide (PI, Becton-Dickinson, San Diego, CA, USA), analysis was carried out according to the manufacturer’s protocol (BMS500F1/100CE Annexin-V-FITC, eBiscience, USA). Finally, the apoptotic cells were counted by FACSscan flow cytometry (Becton Dickinson, Heidelberg, Germany). All experiments were processed independently three times. A minimum of 5 × 10⁵ cell/ml were analyzed for each sample.

RESULTS

Result of determination of IC50 by MTT assay
Cell vitality in the human HCC cell line was analyzed using the MTT assay as described previously. The result of MTT assay indicated that GE inhibits the growth of liver cancer cells significantly in all treatment groups except the control group (P < 0.02) [Figure 1]. The IC50s value for PLC/PRF5 cells were 25 µM of GE at different time periods. According to Figure 1, the percentage of living cells in treatment groups (24, 48, and 72 h) at a concentration of 25 µM were 51, 49, and 47%, respectively (P < 0.001). The effect of GE was dose- and time-dependent. This experiment was repeated three times for each group.

Result of determination of cell viability by MTT assay
The cell vitality in the cells treated with GE at a concentration of 25 µM in different time periods was analyzed using the MTT assay. The amounts of reduced MTT in the all groups treated with GE were significantly lower than that of the control group (P < 0.001) and also in the 72 h treatment group than that of the other experiment groups (P < 0.001) but there isn’t any significant difference between 48 and 72 h treatment groups (P < 0.25). The percentage of living cells in treatment groups (24, 48, and 72 h) were 53%, 48%, 47%, respectively, at a concentration of 25 µM of GE. This experiment was repeated three times for each group [Figure 2].
Result of determination of apoptotic cells by flow cytometry

The cells were treated with 25 µM concentration of GE for different times (24, 48 and 72 h). Flow cytometry was performed to observe the apoptotic cells which had been visualized using Annexin V-FITC and/or PI staining. Flow cytometry analysis indicated that GE at 25 µM concentration induces apoptosis in hepatocellular cancer cells in a time-dependent manner ($P < 0.001$). The amount of apoptotic cells was significantly increased in all three groups, but an apoptotic cell in the 72-h treatment group was more significant [Figure 3]. Percentage of apoptotic cells at different time periods (24, 48, and 72 h) were 44, 56, and 60%, respectively. Apoptotic effects were not observed in DMSO group.

DISCUSSION

Hepatocellular carcinoma is the most common primary malignancy of the liver and the eighth human cancer. The disease is a major complication of liver cirrhosis, and its growth rate is related to the prevalence of risk factors for chronic liver disease. In general, this disease due to a delay in diagnosis has weak pathogenesis.$[1-3]$ The disease...
involves different molecular pathways that can affect the biological and etiologic behavior of the tumor.\(^{[4]}\)

Genistein is a plant-derived estrogen-like compound. The precursor of this bioactive component derived from soy products, grains, nuts, and berries which becomes hormone-like compounds with estrogenic activity by intestinal bacteria.\(^{[42]}\) The plant-derived estrogens have similar activity to beta-estradiol. It is well known that estrogen is involved in various types of cancer such as ovarian, prostate, breast, endometrial and colorectal cancers through different mechanisms that central to these mechanisms is the estrogen receptor (ER) to which estrogen bind.\(^{[49]}\)

Our study clearly indicated that GE has a significant inhibitory effect on the growth of liver cancer cells and induces apoptosis in this cell line with a dose- and time-dependent manner.

Similarly, the same conclusion was reached by other researchers in other cancers; Chang KL \textit{et al.}, reported that GE inhibits proliferation and induces apoptosis in human prostate cancer\(^{[44]}\) and also isolavones inhibit growth of HT-29 and Colo320 cell. Furthermore, other investigators reported that GE inhibits the growth of HCT 116 cells with a dose-dependent manner. This compound inhibits the growth of breast cancer cell lines ADM/MB231, MCF-7 and HBL-100 too.\(^{[45]}\) Peterson and Barnes reported that high concentration of GE (50 or 100 µM) inhibits ER-positive breast cancer cell growth in the human (Peterson and Barnes, 1991, 1996; Pagliacci \textit{et al}. 1994; Chen \textit{et al}. 2003). Many studies have shown that GE induces apoptosis at 50-100 µM concentrations (Pagliacci \textit{et al}. 1994; 2003).

Lee \textit{et al}. reported that GE can inhibit metastasis of cancer cells (Lee JY \textit{et al}. 2012).\(^{[41]}\) On the other hand, Zhang \textit{et al}. and Chen \textit{et al}. reported that consumption of foods rich in soy can reduce the incidence of ovarian cancer. The cancer rates are lower in the women with a high intake of GE (Zhang \textit{et al}. 2004) which is agreement with the Myung’s research that reported that Phytoestrogens reduce the risk of hormone-dependent tumors (Myung \textit{et al}. 2009). Several other studies have demonstrated the inhibitory effect of GE on prostate cancer cells.\(^{[44,46,47]}\) This compound with a concentration of 25 µM or higher concentration inhibits prostatic cancer cells.

These findings confirm our data obviously, but many studies have reported proliferative effects of GE that is opposite of our finding; it has been reported that GE induces apoptosis in the prostatic cancer LAPC-4 cells but has biphasic effect in the LNCap cell line. In fact, this compound has proliferative effect with physiological concentration (<10 µM) and inhibitory response with high concentration (25 or more than 25 µM).\(^{[48]}\)

Similar studies also have shown that a low dose of GE (3.7 µM) has a proliferative effect on human intestinal cells and inhibitory effect with 26-111 µM concentration. Moreover, it stimulates cell growth with <3.7 µM concentration in IEC18 cell line\(^{[59]}\) and also stimulates cell growth in the ER-positive MCF-7 breast carcinoma cells with concentrations of 1 nM to 10 µM/L.\(^{[49]}\)

It has demonstrated that estrogen has proliferative effect on human WRO, FRO, and ARO thyroid carcinoma cells and induces breast and uterus cancer development.\(^{[43,45]}\) Collectively, GE induces significant growth with a low concentration (1 µM) and significant inhibition with a high concentration (25-100 µM). With regard to the estrogenic effect of GE, this compound probably acts through epigenetic mechanism although the mechanism is not fully understood.

The mechanism of estrogenic effects of GE is not fully understood although many pathways have been reported; Kim EJ \textit{et al}. reported that GE inhibits proliferation and induces apoptosis in colon cancer cells (HT-29) by inhibition of insulin-like growth factor-1 receptor and the PI3k/AKT pathway\(^{[60]}\) also including the involvement of the MAPK pathway\(^{[51]}\) which includes ERK-1/2 and AKT as key regulators.\(^{[42]}\) It should be noted that isolavones inhibit the growth of HT-29 and Colo320 cell at the G2/M phase via this mechanism.\(^{[53]}\) Other investigators reported that GE acts through ER (Setchell \textit{et al}. 2005; Setchell and Clerici 2010, Barnes \textit{et al}. 1996).\(^{[44]}\) Chen \textit{et al}. have shown that GE inhibits growth via ER which in response to 50-100 µM concentrations stops cell growth in the G2/M phase (Chen \textit{et al}. 2003). This means that high concentrations of GE (>25 µM) inhibits ER-dependent breast cancer cell growth. Similarly, Pike \textit{et al}. also reported that phytoestrogens act as natural selective ER modulators, depending on the tissue and the presence of coregulator proteins (Pike \textit{et al}. 1999). Besides, many studies have revealed that GE inhibits growth and proliferation of P47D breast cancer cells through signaling molecules associated with ER (ERK1/2, p90RSK, JNK, Akt, and NFκB) depending on the concentration.\(^{[55]}\) Other investigators have reported that 90% of ovarian cancers arise from the epithelial layer which express ERα (Lee JY \textit{et al}. 2012).\(^{[13]}\) In ovarian cancer, GE causes the cell cycle arrest at the G2/M phase. Some researchers have reported the relationship between GE and Bel-2 family (Banerjee \textit{et al}. 2008; Kyle \textit{et al}. 1997; Spinozzi \textit{et al}. 1994). This family is the most well-known apoptosis proteins which divided into apoptotic (Bel-2 and Bel-2) and pro-apoptotic (Bax, Bak, and Bad) groups. GE also induces apoptosis by inhibiting the NFκB and Akt signaling pathways (Brunet \textit{et al}. 1999; Cardone \textit{et al}. 1998; Van Antwerp \textit{et al}. 1996; Wu \textit{et al}. 1996). Furthermore, tyrosine kinase inhibitor is one of the other mechanisms by which GE has an inhibitory effect on the growth of prostate cancer cells via ER.
In summary, the research about effects of GE on HCC is rare, a recent study has shown that GE (10 µM) induces apoptosis in Hep3B cells were cultured in serum-free medium for 6 or 24 h but there is not any report about PLC/PRF5 cell, therefore, we selected this cell line and treated with GE (25 µM). However, based on data from many investigators, as mentioned above, GE is a plant-derived estrogen-like compound with estrogenic activity and biphasic effects that act through ER. [56] It is important to note that GE had an apoptotic effect based on our data while other studies have reported that its effect is biphasic (inhibitory and stimulatory effects). ER acts as a tumor suppressor gene, and GE can increase expression of ER by molecular mechanisms related to apoptosis probably epigenetic pathways are involved in this process which need more researches. We did not perform enzyme activity assay related to methylation and histone modifications and also enzyme immunoassay related to protein levels, but we will perform in next researches and also further researches are needed to determine the clinical applications of GE.

CONCLUSIONS

Our findings suggest that GE may be a potent antiestrogenic compound and can effectively inhibit growth and induce apoptosis in PLC/PRF5 HCC cells. In future studies, the mechanisms and pathways of antiestrogenic effects of GE should be evaluated.

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