An in vitro Evaluation of Apigenin and Apigenin-7-O-glucoside Against HeLa Human Cervical Cancer Cell Line

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Apigenin (API) is a phytocompound belonging to the subclass of flavones that can be found in both functional foods as well as medicinal plants. Recent studies have assigned API antioxidant, anti-inflammatory, anti-spasmodic, anti-viral, anti-thrombotic, anti-angiogenic and chemopreventive potential in vitro on various cell lines and/or in experimental animal models. Apigenin-7-O-glucoside (API-7) is one of its main glycosides and can be commonly found in chamomile flowers, parsley, celery. The aim of this study was to evaluate the in vitro anti-proliferative and pro-apoptotic effects of API and its glucoside (apigenin-7-O-glucoside) against HeLa human cervical cancer cells. Results have shown that in the set experimental conditions both the aglycone as well as the heteroside elicit anti-proliferative and pro-apoptotic potential against the screened cell line, the aglycone being more active than the heteroside.

Keywords: apigenin, apigenin-7-O-glucoside, HeLa cells, proliferation, apoptosis

During history natural products from medicinal plants have played a very important role in protection of human health. The latest studies on this field have exhaustively demonstrated that an impressive number of representatives of Plant Kingdom have therapeutically valuable metabolites; hence, natural products obtain extensive importance to be used for medicinal purposes [1, 2]. In the latest years renewed interest in drugs of natural origins can be clearly observed. The qualities that recommend such an approach relief on some main advantages compared to chemically synthesized molecules that include easy availability and low toxicity at recommended therapeutic doses. Moreover, the easy availability and the fact that plant extracts contain different types of phytochemicals which can act by synergy in order to prevent or cure human disease are other arguments that come to support this kind of approach [3].

Flavonoids (2-phenyl-1, 4-benzopyrone) (C_{15}H_{10}O_{5}) have a fifteen carbon skeleton chemical structure which contain two phenyl rings and a heterocyclic ring. Flavonoids can be classified into six subclasses based on their structural complexity: flavones (artichokes, oranges, lime, grapefruit); flavan-3-ols (goji berries, cranberries, red onion, radish, buckwheat); flavanols (apple, peach, pecan nuts, black berries, dark chocolate); anthocyanins (raspberries, strawberries, black currant, red cabbage); flavanones (red grapes, chicory, green pepper, lemons); and isoflavones (soybean) [4]. Flavonoids are widely distributed in many medicinal plants like buckwheat, Japanese acacia, celery, parsley, chamomile, common hawthorn, ginkgo biloba, soybean, corn, silver birch [5,6]. API, chemically know as 4', 5, 7-trihydroxyflavone, belongs to the flavone subclass of flavonoids. It is the aglycone of some naturally occurring glycosides like apigetrin (API-7), vitexin (apigenin-7-C-glucoside) and isovitexin (apigenin-6-C-glucosid). Apigetrin, also known as cosmosiin is a natural phytoconstituent with better solubility compared to API [7]. API and cosmosiin can be found in many kinds of fruits, vegetables and nutraceuticals, like Apium graveolens L., Pimpinella anisum L., Chrysanthemum morifolium L., Origanum vulgare L., Matricaria chamomilla L., Mentha citrate L., some Rosa L. species.

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Experimental part
Materials and methods
API and API-7 ≥99% (HPLC) were acquired from Sigma-Aldrich, Germany

Cell culture and preparation
The HeLa human cervical cancer cell line (ECACC, European Collection of Cell Cultures, Salisbury, UK) was cultured into Dulbecco’s Modified Eagle’s Medium (DMEM; Gibco BRL, Invitrogen, Carlsbad, CA, USA) supplemented with 10% heat-inactivated fetal calf serum (FCS; PromoCell, Heidelberg, Germany) and 1% penicillin/streptomycin mixture (Pen/Strep, 10,000 IU/mL; PromoCell, Heidelberg, Germany). Experiments were conducted when cells were in the near-confluent phase of growth.

Evaluation of the antiproliferative activity
The growth-inhibitory activity of API and API-7 was tested by standard MTT dye uptake assay on HeLa human cervical cancer cell line as previously described [25]. Briefly HeLa cells were counted and plated at a number of 5000 cells/well into 96-well plates and incubated with the following concentrations of API, respectively API-7: 0.3 µM, 1 µM, 3 µM, 10 µM, 30 µM for 72 h. Afterwards 5 mg/mL MTT (3-(4,5-dimethylthiazolyl-2-yl)-2,5-diphenyltetrazolium bromide) solution was added and incubation was performed for another 4 h. Dimethyl sulfoxide was used in order to dissolve the precipitated formazan crystals and the absorbance was spectrophotometrically measured at 545 nm. Wells with untreated cells, respectively dimethyl sulfoxide were utilized as control.

Evaluation of the pro-apoptotic activity
For the evaluation of the pro-apoptotic activity API, respectively API-7 on HeLa human cervical cancer cell line, DAPI (4’,6-diamidino-2-phenylindole) nucleic acid staining as well as the double Annexin V-FITC staining were performed as previously described [26, 27]. Based on the evaluation of the antiproliferative activity experiments the chosen concentrations of tested agents were 10 µM and 30 µM. Briley, for DAPI staining HeLa cells were cultured in a 6-well plate and stimulated with the chosen concentrations for 72 h. At the end of this time point cells were washed twice with ice-cold PBS. The next step was fixation in 4% paraformaldehyde in PBS for a period of 30 min. The fixation was followed by another wash in
PBS and afterwards permeabilization was induced for 30 min with 2% Triton X-100 (Sigma Aldrich). Cells were blocked afterwards for one hour with 30% FCS in 0.01% Triton X-100. The DAPI staining was realized in the dark for 15 minutes. Visualization of cancer cell nuclei was performed with the help of a fluorescence Olympus IX73 microscope equipped with an integrated DP74 camera (Olympus, Tokyo, Japan). Magnification was 40x. Representative pictures are presented.

Translocation of the phosphatidylserine residues of the HeLa human cervical cancer cell line, as well as alteration of cell membrane and nuclei was detected by Annexin V-FITC combined with a propidium iodide (PI) kit (Invitrogen, ThermoFisher, Vienna, Austria). The number of cells employed in this experiment was 10^6 and the staining steps followed manufacturer’s protocol indications. The cells were analyzed by flow cytometry on FACS Canto II (BD Biosciences, Heidelberg, Germany) using a FACS DIVA device. The experiment was repeated three times. Representative dot-plots are presented.

**Results and discussions**

Results show that both API, the aglycone as well as the API-7 heteroside present an important antiproliferative activity against HeLa human cervical cancer cell line. The calculated IC_{50} values are 12.08 µM for API respectively 18.28 µM for API-7. As expected the aglycone is more active than its conjugated form, but the conjugation element, glucose does not significantly influence the antiproliferative activity. At the highest tested concentration, namely 30 µM an inhibition of proliferation of 81.91± 0.59 for API, respectively 63.97±2.26 for API-7 could be detected. Results that depict logarithmic dose-response curves employing the following concentrations: 0.3; 1; 3; 10; 30 µM can be seen in Figure 1.

![Fig. 1 Antiproliferative activity of API and API-7 against HeLa human cervical cancer cell line](https://revistadechimie.ro)

In order to primary evaluate also a possible apoptotic potential as a secondary mechanism for the anti-cancer activity, DAPI staining, a well-known technique that stains cell nuclei in blue was performed. Incubation with this phytocompound led to a reduction in cell number, as it can be seen by decreased number of stained nuclei in a selected field. Also condensed nuclei as a sign of loss of membrane integrity, condensed chromate filaments or nuclear fragmentation following incubation with API, respectively API-7 could be detected. Representative pictures can be seen in Figure 2.

![Fig. 2 DAPI staining of HeLa human cervical cancer cell line after incubation with API and API-7](https://revistadechimie.ro)
The Annexin V-PI double staining show that HeLa human cervical cancer cells suffer phenomena of early apoptosis, late apoptosis and as well, in a low percentage necrosis. The number of total apoptotic events is linear with the concentration, API being more active than API-7. The most frequent recorded phenomena were the early apoptotic ones. The average value of viable cells following incubation with API -10 μM is 79±2; API -7-10 μM is 84 ±3; API -30 is 67.5±2.5 μM and API -7-30 μM is 72.5 ±2.5. Results can be seen in Figure 3.

Over time, API has gained much popularity due to low toxicity on normal cells compared to cancer cells, this compound proved to possess chemopreventive and therapeutic potential against some variety of cell lines [28].

Human cervical cancer has two major histologic subtypes: squamous cell carcinoma (SCC) - SiHa and adenocarcinoma (AC) – HeLa. Patients with adenocarcinoma have a worse prognosis than patients with squamous cell carcinoma, requiring a specific therapeutic strategy to the cervical cancer subtype [29].

Our results demonstrated that both API, the aglycone as well as the API-7, the heteroside present a moderate antiproliferative and pro-apoptotic activity against HeLa human cervical cancer cell line but in a different manner, the aglycone being more active than the heteroside.

In a similar research line, Pei-Ming Yang et al. analyzed by the consecrated MTT assay the cytotoxicity of API using the human cervical cancer cell lines HeLa and SiHa in a dose range of 5-15 μM. Thus, in response to the treatment, results have shown that HeLa cells were more sensitive than SiHa cells, concluding that the anticancer effect of API may depend on the subtype of cancer of the cervix. A therapeutic approach that involves the combination of IFNγ with API has shown that API has a synergistic effect in combination with IFNγ. To analyze the combination effect of API and IFNγ, it was calculated the combination index (CI). The CI values in HeLa cells were lower than 1, thus it was observed that the combination of IFNγ with API has diminished the cell viability in HeLa cells [29].

In a related approach the effects of API and API-7 on other cell lines have also been studied.

The group of Haihua Bai et al. evaluated the effects of API on MCF-7 human breast cancer cell lines and MCF-10A human normal mammary cells. Both viability as well as apoptosis had a direct proportional dose-dependent and time relationship with API concentration. IC50 was 109.3 ± 3.7 μmol / L following 24 h of incubation and 65.8 ± 4.3 μmol / L following 24 h of incubation of MCF-7 cell lines. Results showed that API inhibits the growth of MCF-7 cells dose-dependent, but without cytotoxic effects on normal breast cells. Cell apoptosis was determined by Annexin V-FITC-PI double staining. It was observed that 24 h exposure to 100 μmol / L API induced a significant augmentation of late apoptotic and necrotic events of MCF-7 cells (from 1.6% to 17.2%). Decreasing the dose to 80 μmol / L, the ratio of early apoptosis in MCF-7 cells changed (from 1.1% to 18.6%). The study concluded that API can induce both apoptosis as well as necrosis in MCF-7 cancer cells [30].

Nouha Nasr Bouzaïene et al. have shown that incubation of B16F10 mouse melanoma cells with apigenin-7-glucoside in the dose range [10-50 μM] using as time points 24 and 48 h inhibited cell proliferation in a dose-dependent manner. To explicate the death mechanism of B16F10 cells induced by apigenin-7-glucoside, it was examined whether API-7 can induce apoptosis by observing morphological changes, these being analyzed and quantified by fluorescent microscopy. At the highest concentrations, it was discovered that the percentage of apoptotic cells increased from 5% (control) to 43% in case of cells incubated for 48 h with API-7 [31].
In the study designed by Whasun Lim et al., the viability of two types of choriocarcinoma cells (JAR and JEG3) was examined after incubation with API at doses of 0.5; 10; 20; 50 and 100 μM. Between concentrations of [10-100 μM], API reduced survival of JEG3 cells. The populations of apoptotic choriocarcinoma cells were estimated by annexin V and propidium iodide (PI) staining. The percentage of apoptotic JAR and JEG3 cells increased following incubation with 20 μM API with the following percentages: 30% for JAR cells and 50% for JEG3 cells [32].

In a similar approach, Shuhua Shan et al. showed a reduction in the survival rate of three colon cancer cell lines (HCT116, HT29 and DLD1) following treatment with API. The IC50 values against HCT116, HT29, and DLD1 cells were 27.9±2.45, 48.2±3.01 and 89.5±8.9 μM [33].

As for an in vivo approach Hui-Hui Cao et al. demonstrated the anti-metastatic effect of API in an animal model of melanoma that involved the injection of B16F10 melanoma cells into syngenic C57BL / 6 mice. Some of these species were treated with API (150 mg / kg) and others with vehicle (0.5% CMC-Na solution). The metastatic nodules in case of API treated mice was significantly reduced compared to those treated with vehicle [34].

Conclusions

In the set experimental conditions, API as well as API-7 presents in vitro antiproliferative and pro-apoptotic potential against HeLa human cervical cancer cell line, the aglycone being more active than the heteroside.

References


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