COMPARATIVE ANALYSIS OF THE ANTI-PROLIFERATIVE EFFECT OF NATURAL PRODUCTS CATECHIN HYDRATE AND EPIGALLOCATECHIN (EXTRACT) APPLIED ON LEUKEMIA LYMPHOCYTES

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Abstract. Cancer diseases are a problem with worldwide importance. However, the lack of selectivity and induction of toxic side effect during conventional cancer therapy continue to provoke the search for innovative approaches. Recent scientific results have reported for synergistic effect between combination of natural products and chemotherapeutic drugs. In this aspect, flavonoids, which are widely distributed in nature, are well known to exhibit numerous biological activities, including antioxidant, antibacterial, anti-inflammatory, anti-viral and anti-cancer effects and may also, play a role in cancer prevention. In the present study, the effects of low concentrations of catechin hydrate and epigallocatechin, Acacia Catechu spray-dried extract, on cell viability of leukemia lymphocytes were investigated and compared, in order to provide an experimental basis for their future incorporation into newly-synthesized biopolymer particles.

Keywords: Cancer, natural products, epigallocatechin, catechin hydrate, cell viability

1. INTRODUCTION

In recent years, the goal in cancer therapy is to achieve selectivity between damaged and healthy cells. The fact that the toxic side effect of applied conventional anticancer drugs is due to induction of oxidative stress and disruption of redox homeostasis in normal (healthy) cells and tissues [1], [2], provoke researchers to search for a new flexible approach in cancer therapy. Our previous results have shown that the combination of natural substances, most of them with redox modulation properties, with conventional or new generation anticancer drugs have shown synergistic cytotoxic effect specific to cancer cells and decrease of oxidative stress in normal cells [3-6]. Flavonoids, a group of natural substances with various phenolic structures, have attracted considerable scientific and therapeutic interest. These compounds are widely distributed in nature and have diverse multiple biological activities, including antioxidant, antimicrobial, anti-inflammatory, antiviral, anti-allergic and anticancer properties [7,8]. Different epidemiological studies have demonstrated decreased risk of cancer diseases, after consumption of foods and drinks, which contain flavonoids (vegetables, fruits and tea) [3,9,10]. The natural products catechin and epigallocatechin, which belong to the flavonoids group, have the ability to scavenge free radicals, reduce the rate of LDL oxidation, inhibit lipid peroxidation and participate in modulation of the immune response in several biological systems [11-14]. In the present study, the effects of Acacia Catechu spray-dried epigallocatechin extract and catechin hydrate, applied in low concentrations (μmol/L), on cell viability of cancer cell line were investigated and compared. The aim is to provide an experimental basis for their future incorporation into chitosan carriers, as well as experimental results for their possible clinical application in cancer therapy.

2. MATERIALS AND METHODS

2.1. Materials

The pure substance catechin hydrate was purchased from Sigma-Aldrich, Steinheim, Germany and the natural product - powdered epigallocatechin extract – from Acacia Catechu was supplied from North India. The experiments were performed on leukemia lymphocytes (Jurkat; RIKEN Bioresource Center, Saitama, Japan) derived from patients with acute lymphoblastic leukemia.

2.2. Cells and treatment protocol

The cells were cultured in RPMI-1640 medium (Sigma-Aldrich, Steinheim, Germany), supplemented with 10% heat-inactivated fetal bovine serum (Gibco, Auckland, New Zealand) and antibiotics (100 U/ml penicillin and 100 μg/ml streptomycin) (Gibco) in a humidified atmosphere at 37 °C with 5% CO2. All cells were collected by centrifugation (1000 × g for 10 min) and replaced in a fresh medium without antibiotics, before treatment. The studied compounds catechin hydrate and epigallocatechin (extract) were dissolved in dimethyl sulfoxide (DMSO; suitable for cell cultures; Sigma-Aldrich). The final concentration of DMSO in

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the cell suspension did not exceed 1%. At this concentration, DMSO did not influence cell viability. The cells (1×10⁶ cells/ml) were incubated with catechin hydrate and epigallocatechin (extract) at the following concentrations: 0.5 μM, 1 μM, 10 μM, 20 μM and 50 μM.

2.3. Cell proliferation and viability assays

MTT assay (Sigma-Aldrich) was used for detection of cell viability and proliferation activity. 50 μL of MTT (3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide) was added to 50 μL cells in fresh medium (place in 96-well plates) and incubated at 37 °C for 1 hour. After incubation, 150 μL of MTT solvent was added to each well and incubated for 15 min in the dark with shaking to fully dissolve MTT formazan. The absorbance at 590 nm was recorded immediately after that, using a microplate reader (TECAN Infinite® M1000, Austria).

2.4. Statistical analysis

All results are expressed as mean ± standard deviation from two independent experiments with three parallel samples for each experiment (n=6). Comparisons between the groups were performed using Student’s t-test. A value of p < 0.05 was considered as significant.

3. Results and Discussion

3.1. Determination of anti-proliferative activity of catechin hydrate and epigallocatechin (extract)

After incubation (24, 48 and 72 hours) with different concentrations of catechin hydrate or epigallocatechin (extract), the cell viability of Jurkat cancer cell line was measured using MTT assay. The results of the effect of catechin hydrate or epigallocatechin (extract), in low concentration ranges (from 0.5 μM to 50 μM ) on the proliferative activity of leukemia lymphocytes, were presented in Figures 1 and 2. The effect of each concentration of the studied compounds was calculated as a percentage of control (untreated cells), where the proliferation activity of the cells was considered 100%. The data demonstrated slight decrease of cell viability of leukemia lymphocytes (up to 20%) after treatment with the studied compounds. Relatively identical effect of both natural compounds applied at all concentrations, after 48 hours incubation was observed. Enhancement of cell viability of Jurkat cells after 24- and 72-hours incubation with epigallocatechin (extract) applied at 20 μM and 50 μM concentrations, was registered. Whilst decreased proliferative activity in the case of cells treated with catechin hydrate at the stated working conditions was observed.

Up-to-date scientific studies also confirm the anti-proliferative activity of catechins. Rayen et al. (2017) investigated a wide concentration range of catechins (from 10 nM to 100 μM) and their results indicated significant reduction of MTT by catechin at concentration of 100 μM. It was also established that epigallocatechin-3-gallate exhibited significantly higher cytotoxic activity at a dose ≥ 10 μM as compared to other studied compounds at these conditions [18].

The experimental evidence, presented by Kirbitz et al. (2011) demonstrated that epigallocatechin gallate, catechin gallate and epicatechin gallate, which are minor components of green tea, inhibited proliferative activity of human pancreatic ductal adenocarcinoma (PDAC) cells in a dose- and time-dependent manner [15]. On the other side, there is data which indicated that applied in high concentrations (200 μM and 400 μM) on non-cancer lymphoid cell line, flavonoids did not modify cell viability in both resting and stimulated lymphocytes after a 24 h incubation period [16].

![Figure 1. Anti-proliferative effect of catechin hydrate on Jurkat cancer cell line at different incubation time.](image1)

![Figure 2. Anti-proliferative effect of epigallocatechin extracted from Acacia Catechu on Jurkat cancer cell line at different incubation time.](image2)
The similarity of the anti-proliferative effect of catechin hydrate and epigallocatechin (extract) could be due to the same molecular mechanism of action of the studied compounds. It is proposed that the anti-inflammatory and immunomodulatory activity of flavonoids is due to regulation of NF-κB signaling (nuclear factor kappa-light-chain-enhancer of activated B cells) [11]. In this aspect, Mackenzie et al. (2004) established that flavonoids, including catechin, were implicated in the modulation of phorbol 12-myristate 13-acetate (PMA)-induced NF-κB activation in Jurkat T cells by the regulation of IL-2 at the level of transcription. The obtained results displayed that preincubation for 24 hours with catechin, applied in different concentration range (from 1.7 μM to 17.2 μM), decreased PMA-induced NF-κB binding activity, through a direct interaction of flavonoids with NF-κB proteins, inhibition of the binding of active NF-κB to κB sites and the transactivation of the NF-κB-driven gene IL-2 [11]. Thus, it has been considered that the intracellular mechanism of action of catechins is associated with an influence on the immune response by modulating NF-κB activation. In this aspect, the study of Suhaie et al. (2019) presented a corroboration of the impact of flavonoids on the specific for cancer cells NF-κB molecular pathway. The provided results form molecular docking demonstrated that epicatechin, epigallocatechin, epicatechin gallate and epigallocatechin gallate block NF-κB binding site through interaction with its different amino acids residues. The scientific team reported the formation of protein-ligand interaction between NF-κB and the studied flavonoids [17]. An earlier study demonstrated that catechin gallate and epicatechin gallate inhibited the proliferation of human pancreatic ductal adenocarcinoma (PDAC) cells due to influence of modulation on NF-κB activity. In the same report, the evidences outlined that epicatechin gallate inhibited TNFα-induced activation of NF-κB, the consequent secretion of pro-inflammatory and invasion of promoting proteins such as IL-8 and the urokinase plasminogen activator (uPA). In this regard, due to the observed anti-inflammatory potential of flavonoids, especially of epicatechin gallate, they could be applied as supplements to anticancer therapies [15].

4. CONCLUSION

In summary, the results of the present study provide evidence that catechin hydrate and epigallocatechin, extracted from Acacia Catechu, applied at low concentrations (μmol/L), displayed slight decrease of cell viability of leukemia lymphocytes. Thus, future studies of the potential of chitosan/epigallocatechin-loaded chitosan-based carriers, in combination with chemotherapeutic drugs could present essential data for the probable benefits of their application in cancer therapy. However, the in vitro anticancer and anti-inflammatory activity mechanisms of such bioflavonoid-biopolymer carrier formulations on different cancer cell lines need to be precisely investigated.

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REFERENCES


Figure 3. Comparative analysis of anti-proliferative activity of catechin hydrate (in blue) and epigallocatechin extracted from Acacia Catechu (in yellow) on Jurkat cell line after 48 hours incubation time. The grey color indicates untreated cells (control). (Mean±SD, n=6). Two levels of significances were considered: p<0.05 (*) and p<0.01 (**).

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