SYNERGISTIC EFFECT OF QUERCETIN AND EPIGALLOCATECHIN-3-GALLATE AS AGENTS TO PREVENT CONNECTIVE TISSUE DISINTEGRATION IN THE PERIODONTIUM OF RATS UNDER SYSTEMIC AND LOCAL ADMINISTRATION OF LIPOPOLISACCHARIDE OF SALMONELLA TYPHI*

Yelins’ka A.M., Kostenko V.O.

Ukrainian Medical Stomatological Academy, Poltava, Ukraine

The aim of the present study was to investigate the co-effect produced by water-soluble form of quercetin and epigallocatechin-3-gallate (EGCG) on biochemical markers of periodontal organic matrix depolymerization under systemic administration and local application of S. typhi lipopolisaccharide (LPS). The studies were conducted on 30 white rats of the Wistar line weighing 180-220 g, divided into 5 groups: the 1st included intact animals, the 2nd was made up of animals after the combined systemic and local LPS administration, the 3rd and 4th groups included animals, which were being given injections with water-soluble form of quercetin (10 mg / kg) and EGCG (21.1 mg / kg) respectively 3 times a week, starting on the 30th day of the systemic LPS administration, and the 5th group involved rats, which were injected with co-administered water-soluble form of quercetin and EGCG. It has been found out that the co-effect produced by quercetin and EGCG under systemic and local LPS administration is accompanied with reduced concentration of N-acetylserymamic acid (NANA) by 31.8 and 32.8% respectively in the soft periodontal tissues. At the same time combined use of quercetin and EGCG under experimental conditions led to the decrease in the FHP content in the alveolar bone by 24.5 and 20.2% respectively compared with values for the animals received separate quercetin and EGCG during the experiment. However, no differences have been detected between the groups exposed to combined or separate action of the above mentioned agents in the experiment when analyzing free hydroxyproline (FHP) and glycosaminoglycans (GAGs) content in the soft tissues of periodontium. It has been found out that the co-administration of water-soluble form of quercetin and epigallocatechin-3-gallate under systemic and local introducing of S. typhi lipopolysaccharide has been proven to be more effective means for preventing and correcting periodontal connective tissue disruption than this occurs at separate administration of each of the polyphenols.

Key words: quercetin, epigallocatechin-3-gallate, lipopolisaccharide, connective tissues disintegration, periodontium.

Introduction

Epigallocatechin-3-gallate (EGCG) is a polyphenol found in green tea (Camellia sinensis), that promotes the activation of Nrf2 transcription factor (Nuclear Factor Erythroid 2-Related Factor 2) due to proteolysis of an inhibitory protein Keap1 [4]. This pathway enhances antioxidant activity of a number of enzymes through cis-acting enhancer sequence, known as antioxidant response element (ARE) [10, 13]. In our previous reports we have demonstrated that the administration of EGCG under modeled systemic inflammatory response (SIR) is an effective means of preventing and correcting the disruption of periodontium connective tissue in rats: it reduces collagenolysis and depolymerization of proteoglycans and glycoproteins [17]. A particular role of ARE is determined by the fact that transcriptional nuclear factors

* To cite this English version: Yelins’ka A.M., Kostenko V.O. Synergistic effect of quercetin and epigallocatechin-3-gallate as agents to prevent connective tissue disintegration in the periodontium of rats under systemic and local administration of lipopolisaccharide of salmonella typhi. // The Medical and ecological problems. - 2019. - Vol 23, № 5-6. - P. 42-44.
kB (NF-κB) and activator protein 1 (AP-1) activation depends on its activity [13]. Nowadays, the ability of quercetin to influence on the activity of enzymes involved in the degradation of phospholipids (phospholipase, lipoxgenase, cyclooxygenase), and to block oxidative stress-dependent connective tissue disruption has already been proven [5]. There is evidence suggesting quercetin ability to activate the Nrf2-dependent HO-1 pathway and suppress ubiquitin-dependent proteolysis of NF-κB with an inhibitory protein kB, which impairs the degradation of the latter under the protective action [3].

This creates the prerequisites for eliminating the possibility of NF-κB-dependent expression of numerous genes, a significant number of which encodes hystolic matrix metalloproteinases (MMP), proinflammatory cytokines, inducible NO-synthase, etc [6] Recently, it has been reported that an administration of NF-κB inhibitors is accompanied by an increase in periodontal collagen-protective activity of L-arginine [8] However, the combined use of EGCG and quercetin for prevention and correction of connective tissue disintegration in periodontium of rats under systemic and local pathogens action is still remaining unclear.

The aim of the present study was to investigate the co-effect produced by quercetin and EGCG on biochemical markers of periodontal organic matrix depolymerization under systemic and local administration of S. typhi lipopolysaccharide (LPS).

Materials and methods

The studies were conducted on 30 white rats of the Wistar line weighing 180-220 g, divided into 5 groups: the 1st included intact animals, the 2nd was made up of animals after the combined systemic and local LPS administration, the 3rd and 4th groups included animals, which were given injections with water-soluble form of quercetin and EGCG respectively, and the 5th group involved rats, which were injected with co-administered water-soluble form of quercetin and EGCG. For systemic administration, S. typhi LPS (pyrogenalum, "Medgama", Russia) was injected intraperitoneally in a dose that stimulated rise in temperature by 1.5 °C according to the scheme [19]; during the first week, 4 minimum pyrogenic doses (MPD) of 0.4 µg / kg of body weight were given 3 times a week. During the following 7 weeks of the experiment, rats were given 4 MPD / kg of body weight once a week. For local administration, S. typhi LPS was introduced once in as dose of 1 µg / kg, equally divided into four injections into the gum at the level of the second molars 7 days prior to the decapitation (acute gingivitis model). Water-soluble form of quercetin (corditin, "Borschchak-hivskiy CPP", Ukraine) was administered intraperitoneally in a daily dose of 10 mg / kg once calculated for quercetin [5], and EGCG (Sigma-Aldrich, Inc., USA) was administered in a dose of 21.1 mg / kg [17] 3 times a week, starting on the 30th day of the systemic LPS administration.

The research was guided by the principles of biomedical ethics. The animals were decapitated under etheal anesthesia. The level of collagenolysis was assessed by the content of free hydroxyproline (FHP) [12]. The process of depolymerization of proteoglycans and sialoglycoproteins was evaluated by determining their monomers – glycosaminoglycans (GAGs) [11] and N-acetyleneuraminic acid (NANA) [9] respectively.

The findings obtained were statistically processed. To verify the normality distribution, the calculation of the Shapiro-Wilk criterion was applied. If they corresponded to the normal distribution, then the Student’s t-test was used to compare independent samples. When the results ranges were not subject to normal distribution, statistical processing was performed using a nonparametric method, the Mann-Whitney test. Statistical calculations were performed using the "StatisticaSoft 6.0" program.

Results and discussion

Systemic and local LPS co-administration led to significant changes in the biochemical markers of periodontal organic matrix destruction (Table 1). Thus, the FHP and GAGs content went up by 2.25 times, NANA - by 2.28 times in the soft periodontal tissues. Concentration of these compounds in the calcified components of periodontium (alveolar bone) increased as well: FHP - by 2.45 times, GAGs - by 2.25 times, NANA - by 3.55 times.

Table 1

<table>
<thead>
<tr>
<th>Groups of the animals</th>
<th>Soft components (gingiva and periodontal ligament)</th>
<th>Calcified components (alveolar bone)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>FHP, µmol/g</td>
<td>GAGs, µmol/g</td>
</tr>
<tr>
<td>Control group I (intact animals)</td>
<td>4.08 ±0.48</td>
<td>1.93 ±0.34</td>
</tr>
<tr>
<td>Control group II (combined systemic and local LPS introduction)</td>
<td>9.20 ±0.23</td>
<td>4.36 ±0.17</td>
</tr>
<tr>
<td>+ water-soluble form of quercetin</td>
<td>5.51 ±0.42</td>
<td>2.59 ±0.40</td>
</tr>
<tr>
<td>+ EGCG</td>
<td>5.16 ±0.44</td>
<td>2.27 ±0.30</td>
</tr>
<tr>
<td>+ water-soluble form of quercetin + EGCG</td>
<td>4.04 ±0.42</td>
<td>1.93 ±0.27</td>
</tr>
</tbody>
</table>

Note: * p<0.05 compared with values in the control group I (intact rats); ** with control group II; *** with animals received quercetin only under experiment; **** p<0.05 with rats received EGCG only under experiment.

Recent studies have demonstrated the activation of NF-κB and AP-1 enhances MMP-13 (collagenase-3) gene expression [2]. Introduction of an inhibitor of the nuclear translocation of NF-κB 4-methyl-N-(3-phenylpropyl)
benzene-1,2-diamine in systemic inflammatory response under experimental metabolic syndrome reduces the amount of FHP and GAGs in periodontal tissues [7].

It has been found that the separate administration of both quercetin and EGCG reduces the concentration of FHP by 42.3 and 43.9%, the content of GAGs by 40.6 and 47.9%, and NANA content by 40.2 and 39.3% respectively in the soft periodontal tissues compared with the relevant findings in the 2nd group of the test animals. In this experimental model, the concentration of these compounds in the calcified components of periodontium (alveolar bone) lowered as well: FHP - by 47.8 and 50.6%, GAGs - by 42.2 and 48.2%, NANA - by 54.5 and 49.6% respectively compared with the relevant findings in the 2nd group of the animals.

Co-effect produced by quercetin and EGCG under systemic and local LPS administration demonstrates the reduced NANA concentration by 31.8 and 32.8% respectively in the soft periodontal tissues compared with values for the animals received separate quercetin and EGCG during the experiment. However, no differences were found out between the groups exposed to either combined or separate action of the above mentioned agents in the experiment when assessing FHP and GAGs content in the soft tissues of periodontium.

At the same time the combined use of quercetin and EGCG under experimental condition led to decrease in the FHP content in the alveolar bone by 24.5 and 20.2% respectively compared with values for the animals received separate quercetin and EGCG. NANA concentration was reduced by 35.0 and 41.3% respectively. No differences were found between the groups exposed to either combined or separate action of the agents in the experiment when assessing GAGs content in the calcified components of periodontium.

The results obtained have demonstrated the synergic effect produced by water-soluble form of quercetin and EGCG on the correction of connective tissue disruption in the periodontal tissues.

Recent studies have shown the potential of Keap1 / Nrf2 / ARE system to control other redox-sensitive elements, including NF-κB and AP-1 [14]. We have also found out the activation of NF-κB and AP-1 is an important component in the mechanism of free radical injury and extracellular matrix destruction in the periodontal tissues during systemic inflammatory response [1, 7, 15, 16].

Co-administration of water-soluble form of quercetin and EGCG enables to limit NF-κB and AP-1-dependent mechanisms of periodontal disintegration more effectively as well as to enhance periodontium resistance associated with ARE-dependent gene expression.

Thus, the co-administration of water-soluble form of quercetin and epigallocatechin-3-gallate under systemic and local introduction of S. typhi lipopolysaccharide has been proven to be more effective means for preventing and correcting periodontal connective tissue disintegration than this occurs at separate administration of each of the polyphenols.

References

7. Ljashenko LI, Denisenko SV, Kostenko VA Role of transcription nuclear factor κB in mechanisms of free radical processes impairment and connective tissue disorganization in periodontium under modeled metabolic syndrome. Aktualni problemy suchasnoyi medytsyny: Visn Ukr med stomatol akad. 2014; 14(1):97-100. [In Ukrainian].
11. Sharayev PN Method for the determination of glycosaminoglycans in biological fluids. Lab delo. 1987;(5):530-2. [In Russian].
12. Tetyanets SS. Method for the determination of free hydroxyproline in serum. Lab delo. 1985;(1):61-6. [In Russian].
18. Yelins’ka AM, Shvaykovs’ka OO, Kostenko VO Influence of ammonium pyrrolidine dithiocarbamate on production of reactive oxygen and nitrogen species in tissues of periodontium and salivary glands of rats exposed to systemic administration of Salmonella typhi lipopolysaccharide. Fiziol Zh. 2018;64(5):63-9. [In Ukrainian].

Материал надійшов до редакції 23.08.2019.