Chronic systemic inflammatory response syndrome (SIRS) underlies many diseases (sepsis, atherosclerosis, diabetes mellitus). According to research data of recent years the key role in the development of SIRS is played by the activation of various nuclear transcription factors. The work was aimed at studying the role of such transcription factor as activator protein 1 (AP-1) in the development of oxidative and nitrosative stress in soft periodontal tissues during chronic systemic inflammatory response (SIRS). The experiment was carried out on 24 the Wistar rats. We induced SIRS by bacterial lipopolysaccharide of Salmonella typhi (0.4 μg/kg) intraperitoneal injection. We studied changes in the functioning of the nitric oxide (NO) cycle, the production of superoxide anion radical (O$_2^•$-) and the activity of antioxidant enzymes in soft periodontal tissues homogenate. We used SR11302 as an AP-1 inhibitor (15 mg/kg) for 2 months. We established that during the SIRS modeling, the activity of antioxidant enzymes in soft periodontal tissues decreased with a simultaneous increase in the production of O$_2^•$. SIRS elevated the production of NO by inducible NO-synthase (iNOS) and nitrite reductases. The nonoxidative cleavage of L-arginine under this condition was also increased. The concentration of peroxynitrite (ONOO-) was shown to be elevated more than 2-fold. The inhibition of AP-1 by SR11302 normalized the functional state of the NO cycle, reduced O$_2^•$- production and restored the activity of antioxidant enzymes. In this way, under SIRS conditions, “vicious circle” of ONOO$^-$ formation is formed.

SIRS in soft periodontal tissues poses a threat of oxidative and nitrosative stress development. Usage of AP-1 activation inhibitor SR11302 breaks “vicious circle” of ONOO$^-$ formation.

Key words: periodontium, systemic inflammatory response, activator protein 1 (AP-1), superoxide anion radical, nitric oxide cycle.
role of activator protein-1 (AP-1) in the development of SIRS-induced oxidative and nitrosative stress in periodontal tissues has not been sufficiently studied. The aim of this work is to study the role of AP-1 in the development of oxidative and nitrosative stress in soft periodontal tissues under the SIRS.

**Materials and Methods**

We carried out the study on 24 mature male Wistar rats weighing 200-220 g. Bioethical committee of Ukrainian Medical Stomatological Academy approved all experimental procedures with experimental animals. Animals were divided into 3 groups: control animals (8), SIRS group (8), and animals (8), which received the AP-1 inhibitor, namely (2E,4E,6Z,8Z)-3-methyl-7-(4-methylphenyl)-9-(2,6,6-trimethylcyclohexen-1-yl)nona-2,4,6,8-tetraenoic acid (SR11302) in a dose of 15 mg/kg (iAP-1 group).

The experiment lasted 2 months. The chronic systemic inflammatory response was induced by injection of bacterial lipopolysaccharide of *Salmonella typhi* (Pyrogenalum, Medgamal, Russia) intraperitoneally in a dose of 4 minimal pyrogenic doses (MPD) per 1 kg of animal body weight that corresponds to 0.4 μg/kg. We administered Pyrogenalum three times during the 1st week of the experiment to trigger an acute systemic inflammatory response; then in order to develop and maintain a chronic systemic inflammatory response Pyrogenalum was administered once a week for 1 month [5]. The animals from the SIRS group continued to receive Pyrogenalum once a week until the end of the experiment. After the development of SIRS, the animals from the iAP-1 group received injections of an aqueous solution of SR11302 intraperitoneally twice a week for 2 months. The volume of the injected liquid did not exceed 0.2 ml.

We removed animals from the experiment under ether anesthesia by decapitation. We carried out all biochemical studies in 10% periodontal tissues homogenate.

Determination of total NO-synthases, arginases, nitrate and nitrite reductases activities and concentration of peroxynitrite was performed as described in [6]. In addition to method [6] to determine the activity of constitutive NO-synthases (cNOS) we modified procedure proposed in [6]. The modification consists from adding 0.1 ml of 1% (weight/volume) solution of aminoguanidine hydrochloride (98% Sigma Aldrich) to the first aliquot (0.2 ml of 10% homogenate, 2.5 ml 0.1 M Tris-buffer, 0.3 ml 320 mM L-arginine water solution and 0.1 ml 1 mM NADPH solution) and changing incubation time to 60 min. The rest of the method was carried out as described in [6]. The activity of inducible NO-synthase (EC 1.14.13.39, iNOS) was evaluated by subtracting the cNOS activity from the overall activity of NO-synthases (EC 1.14.13, NOS).

We evaluated the production of the superoxide radical anion (O$_2^-$) by estimation of concentration of diformazan, formed in reaction of O$_2^-$ with nitroblue tetrazolium (IUPAC: 2-[2-methoxy-4-[3-methoxy-4-[3-(4-nitrophenyl)-5-phenyltetrazol-2-ium-2-yl]phenyl]phenyl]-3-(4-nitrophenyl)-5-phenyltetrazol-2-imium) [7]. In order to estimate influence of cellular organelles (mitochondria and endoplasmic reticulum) and tissue phagocytes specific reaction conditions described in [7] were used. The activity of superoxide dismutase (EC 1.15.1.1, SOD) was determined according to the recommendations [8]; the activity of catalase (EC 1.11.1.6, CAT) was assessed by the method [9]. Protein concentration was determined by Biurette method.

The results were analyzed for normality by the Shapiro-Wilk test. Then, non-parametric analysis by Kruskal-Wilks test and comparison of groups by the Mann-Whitney test were carried out. The difference between groups was considered statistically significant when $P < 0.05$. 

**Results and Discussion**

SIRS notably increased production of O$_2^-$ from all its sources in periodontal tissues homogenate (Table 1). Basal production of O$_2^-$ increased by 16.9%. O$_2^-$ production by the mitochondrial electron transport chain (ETC) elevated in 1.47 times. O$_2^-$ production under SIRS conditions by microsomal ETC and NO-synthase increased by 61.6%. Periodontal phagocytes increased O$_2^-$ production by 19.3%.

SIRS was accompanied by decreased activity of antioxidant enzymes. The activity of SOD and CAT was reduced by 32.8 and 9.5% respectively.

The production of nitric oxide (NO) by L-arginine-dependent pathway was elevated by 3.9 times under SIRS conditions (Table 2), activity of nitrite reductases increased by 34.7%, nitrate reductases increased their activity by 55.0%. cNOS activity...
**Table 1.** $O_2^•$- production and antioxidant enzymes activity in rat periodontium under conditions of chronic systemic inflammatory response and usage of AP-1 inhibitor ($M ± m$, $n = 8$)

<table>
<thead>
<tr>
<th>Parameters</th>
<th>Control</th>
<th>SIRS</th>
<th>SIRS + AP-1 inhibitor</th>
</tr>
</thead>
<tbody>
<tr>
<td>Production of $O_2^•$, nmol/s per g of tissue:</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Basic</td>
<td>0.77 ± 0.03</td>
<td>0.90 ± 0.01*</td>
<td>0.81 ± 0.033*</td>
</tr>
<tr>
<td>From microsomal ETC</td>
<td>18.64 ± 0.57</td>
<td>30.13 ± 0.29*</td>
<td>18.87 ± 0.57*</td>
</tr>
<tr>
<td>From mitochondrial ETC</td>
<td>11.73 ± 1.44</td>
<td>28.95 ± 0.46*</td>
<td>11.83 ± 1.42*</td>
</tr>
<tr>
<td>From phagocytes</td>
<td>1.14 ± 0.02</td>
<td>1.36 ± 0.01*</td>
<td>1.17 ± 0.02*</td>
</tr>
<tr>
<td>SOD activity, c.u.</td>
<td>8.32 ± 0.45</td>
<td>5.59 ± 0.56*</td>
<td>8.01 ± 0.54*</td>
</tr>
<tr>
<td>Catalase activity, μkat/g of tissue</td>
<td>0.241 ± 0.006</td>
<td>0.218 ± 0.005*</td>
<td>0.239 ± 0.005*</td>
</tr>
</tbody>
</table>

*Data is significantly different from control group ($P < 0.05$); †data is significantly different from chronic systemic inflammation response group ($P < 0.05$)

**Table 2.** The functioning of the nitric oxide cycle in rats periodontium under conditions of chronic systemic inflammatory response and usage of AP-1 inhibitor ($M ± m$, $n = 8$)

<table>
<thead>
<tr>
<th>Parameters</th>
<th>Control</th>
<th>SIRS</th>
<th>SIRS + AP-1 inhibitor</th>
</tr>
</thead>
<tbody>
<tr>
<td>NOS activity, nmol (NO$_2^•$)/min·mg of protein:</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Total</td>
<td>0.61 ± 0.16</td>
<td>2.39 ± 0.24*</td>
<td>0.61 ± 0.17*</td>
</tr>
<tr>
<td>cNOS</td>
<td>0.38 ± 0.14</td>
<td>0.48 ± 0.06</td>
<td>0.37 ± 0.14</td>
</tr>
<tr>
<td>iNOS</td>
<td>0.25 ± 0.04</td>
<td>1.91 ± 0.19*</td>
<td>0.24 ± 0.04*</td>
</tr>
<tr>
<td>ONOO$^•$ concentration, μmol/g of tissue</td>
<td>0.67 ± 0.20</td>
<td>1.41 ± 0.13*</td>
<td>0.71 ± 0.23*</td>
</tr>
<tr>
<td>Total arginase activity, nmol (L-ornithine)/min·mg of protein</td>
<td>1.05 ± 0.16</td>
<td>4.05 ± 0.27*</td>
<td>1.00 ± 0.18*</td>
</tr>
<tr>
<td>Total nitrate reductase activity, nmol (NO$_3^•$)/min·mg of protein</td>
<td>2.89 ± 0.48</td>
<td>4.48 ± 0.56*</td>
<td>2.74 ± 0.35*</td>
</tr>
<tr>
<td>Total nitrite reductase activity, nmol (NO$_2^•$)/min·mg of protein</td>
<td>2.77 ± 0.35</td>
<td>3.73 ± 0.22*</td>
<td>2.53 ± 0.38*</td>
</tr>
</tbody>
</table>

*Data is significantly different from control group ($P < 0.05$); †data is significantly different from chronic systemic inflammation response group ($P < 0.05$)

did not show statistically significant changes, while the iNOS activity increased by 7.6 times. Arginase activity (EC 3.5.3.1) increased by 3.9 times. SIRS enhances the reduction of nitrates to nitrites by 1.6 times. The concentration of peroxynitrite (ONOO$^•$) was twice as much when compared with the control group.

The inhibition of AP-1 during the SIRS modeling reduced basic $O_2^•$ production by 10%. Production of $O_2^•$ by mitochondrial and microsomal ETC was reduced by 37.4% and 59.1%, respectively. Usage of SR11302 slowed down (by 14.0%) $O_2^•$ production by phagocytes of periodontal tissues. SOD activity under conditions of AP-1 inactivation during SIRS increased by 43.3%. CAT activity increased by 9.6%.

Using SR11302 reduced overall NOS activity by 73.6%. iNOS activity dropped by 87.4%. There were no changes in cNOS activity as compared with the SIRS modeling group. Inhibition of AP-1 reduced the ONOO$^•$ concentration by 2.0 times. Arginases activity decreased by 4.0 times. Nitrate reductases activity decreased by 1.63 times, nitrite reductases activity decreased by 32.17%.

**Mechanisms of oxidative-nitrosative stress development in periodontium during SIRS.** Increased
O$_2^\cdot$ production may be considered as a result of increased activity of arginases, which according to Rabelo L.A. et al., are pro-oxidants [10]. $K_m$ for L-arginine of arginase I and II (approximately 10 mmol/l) [11], which is much higher than $K_m$ for L-arginine of different NOS isoforms (approximately 5 μmol/l for iNOS and 3 μmol/l for eNOS) [12]. Therefore excessive arginase activation can lead to L-arginine deficiency for cNOS and they may shift to O$_2^\cdot$ production instead of NO production. $V_{max}/K_m$ for L-arginine of both iNOS and arginases are close so competition between them affects iNOS less than it affects nNOS or eNOS isoforms. According to the results obtained, cNOS activity does not change during SIRS modeling, thus, an increase in the O$_2^\cdot$ production by microsomal ETC is not associated with the cNOS substrate deficiency. A characteristic feature of our SIRS model is the long-term administration of bacterial lipopolysaccharide that leads to hypercytokinemia (increased IL-1 and TNF-α). Cao Y. reported that stimulation of chondrocytes with pro-inflammatory cytokines (IL-1 and TNF-α) results in an increase in the production of reactive oxygen species (ROS) by cell mitochondria [13]. IL-1β can enhance the activity of microsomal NADPH oxidase and lead to over-production of ROS by the microsomal ETC [14]. Elevated O$_2^\cdot$ production can lead to oxidative stress development in periodontal tissues.

During SIRS modeling in periodontal tissues we observed excessive NO production caused by iNOS activation. Enhanced activity of iNOS may be the result of Pyrogenalum administration. Pyrogenalum, according to its pharmacodynamics, leads to increased concentration of IL-1 in blood, which can result in increased gene expression of iNOS [15]. At the same time we found out an increase in nitrite reductases activity that also yields NO. The elevation of nitrite reductases activity during SIRS conditions can be regarded as a protective mechanism aimed at preventing accumulation of free nitrates in periodontal tissues. Under physiological conditions nitrates get oxidized to nitrites and then are excreted with urine. Increased nitrate reductase activity in soft parodontal tissues makes this way of nitrates excretion difficult. Hence, the development of nitritive stress in the periodontal tissues is quite possible.

Reduction of antioxidant enzymes activity during SIRS modeling may develop as a consequence of depletion of these systems or “interception” of O$_2^\cdot$ by nitrogen oxide with the following ONOO$^\cdot$ formation. The velocity rate of SOD reaction with O$_2^\cdot$ is $\kappa = 2.0 \cdot 10^9$ M$^{-1}$s$^{-1}$, while $\kappa = 6.7 \cdot 10^8$ M$^{-1}$s$^{-1}$ for the O$_2^\cdot$ reaction with NO [16]. Under the conditions of NO hyperproduction in periodontal tissues, a sharp rise in the ONOO$^\cdot$ concentration confirms the possibility of “interception” of O$_2^\cdot$. Increased ONOO$^\cdot$ production can lead to the development of nitrosative stress.

ONOO$^\cdot$ can oxidize tetrahydrobiopterin to its inactive radical leading to NOS uncoupling and subsequent O$_2^\cdot$ production. Increased total activity of NOS indirectly proves NO hyperproduction, while increased nitrite reductases activity limits physiological NO removal. This leads to excessive levels of both NO and O$_2^\cdot$ which in turn leads to ONOO$^\cdot$ formation and creates “vicious circle”.

Peroxynitrite is a powerful oxidant formed in vivo, that can directly react with different biomolecules by one (i.e. reaction with transition metals) or two-electron (i.e. reaction with thiols) oxidations. ONOO$^\cdot$ in biological systems can react with carbon dioxide (CO$_2$) in equilibrium with bicarbonate (HCO$_3^-$) forming ‘NO$_2^-$ and CO$_2^\cdot$ radicals, highly oxidant species that can in turn mediate oxidative damage to biomolecules [17, 18].

**Role of AP-1 inhibition in oxidative-nitrosative stress development in periodontal tissues during SIRS modeling.** Blockade of AP-1 activation reduces the O$_2^\cdot$ production. As a mechanism that underlies this effect, we can consider the decrease in pro-inflammatory cytokines levels, which are controlled by AP-1. Tao X. et al. showed that the AP-1 inhibition by flavonoids from Rosa laevigata is accompanied by a decrease in ROS production and lowering IL-1, IL-6 and TNF-α expression [19]. Kankaanranta H. et al. demonstrated that TNF-α is capable of leading to apoptosis only under the conditions of imbalance between NF-κB and AP-1 [20]. The importance of a balance between transcription factors is also indicated by Fujioka S. et al. They found out that NF-κB activation in response to stimulation by TNF-α, but not by IL-1, is able to inhibit AP-1 activation [21].

Jang B et al. proved that AP-1 is not directly linked to iNOS activation when lipopolysaccharides are administered [22]. However, it has been shown that AP-1 activation leads to suppression of endothelial NOS isoforms (EC 1.14.13.39, eNOS), by enhancing oxidative stress [23]. The lack of eNOS activity and oxidative stress can evoke iNOS activation. Thus we can suggest that AP-1 inhibition reduces iNOS activity by decreasing oxidation load.
and restoring the eNOS activity. In present study we did not evaluate separate activities of nNOS and eNOS, which is limitation of our study. We studied the sum of their total activities (cNOS). The absence of changes in cNOS activity during SIRS modeling can be explained by decrease in eNOS activity with simultaneous increase in nNOS activity, since AP-1 has the capacity to increase nNOS activity [24].

AP-1 controls arginase activation through binding of Fos to the AP-1 site in the arginase 1 promoter [25]. Thus inhibition of AP-1 results in arginase activity decrease, thus lowering competition between NOS and arginases for substrate.

SIRS modeling by the introduction of bacterial lipopolysaccharide (Pyrogenalum) leads to excessive production of O$_2^•$, NO and ONOO$^-$ in soft periodontal tissues. Under SIRS conditions “vicious circle” of ONOO$^-$ formation is formed. SIRS in soft periodontal tissues poses a threat of oxidative and nitrosative stress development.

Usage of AP-1 activation inhibitor SR11302 breaks “vicious circle” of ONOO$^-$ formation.

Conflict of interest. Authors have completed the Unified Conflicts of Interest form at http://ukrbiochemjournal.org/wp-content/uploads/2018/12/coi_disclosure.pdf and declare no conflict of interest.


