Research Article
Nitroso-Oxidative Stress, Acute Phase Response, and Cytogenetic Damage in Wistar Rats Treated with Adrenaline

Milena Radaković,1 Sunčica Borozan,2 Ninoslav Djelić,1 Saša Ivanović,3 Dejana Ćupić Miladinović,3 Marko Ristanić,1 Biljana Spremo-Potparević,4 and Zoran Stanimirović1

1Department of Biology, Faculty of Veterinary Medicine, University of Belgrade, Bulevar oslobodjenja 18, 11000 Belgrade, Serbia
2Department of Chemistry, Faculty of Veterinary Medicine, University of Belgrade, Bulevar oslobodjenja 18, 11000 Belgrade, Serbia
3Department of Pharmacology and Toxicology, Faculty of Veterinary Medicine, University of Belgrade, Bulevar oslobodjenja 18, 11000 Belgrade, Serbia
4Department of Physiology, Faculty of Pharmacy, University of Belgrade, Vojvode Stepe 450, 1221 Belgrade, Serbia

Correspondence should be addressed to Milena Radaković; mradakovic@vet.bg.ac.rs

Received 30 March 2018; Revised 19 July 2018; Accepted 12 September 2018; Published 21 November 2018

Guest Editor: Mohamed M. Abdel-Daim

Copyright © 2018 Milena Radaković et al. This is an open access article distributed under the Creative Commons Attribution License, which permits unrestricted use, distribution, and reproduction in any medium, provided the original work is properly cited.

This study is aimed at analysing biochemical and genetic endpoints of toxic effects after administration of adrenaline. For this purpose, the study was carried out on Wistar rats and three doses of adrenaline were used: 0.75 mg/kg, 1.5 mg/kg, and 3 mg/kg body weight. To achieve these aims, we investigated the effects of adrenaline on catalase (CAT), Cu, Zn-superoxide dismutase (SOD), malondialdehyde (MDA), nitrite (NO₂⁻), carbonyl groups (PCC), and nitrotyrosine (3-NT). Total activity of lactate dehydrogenase (LDH), its relative distribution (LDH₁–LDH₅) activity, level of acute phase proteins (APPs), and genotoxic effect were also evaluated. The obtained results revealed that all doses of adrenaline induced a significant rise in CAT activity, MDA level, PCC, NO₂⁻, and 3-NT and a significant decrease in SOD activity compared to control. Adrenaline exerted an increase in total activity of LDH, LDH₁, and LDH₂ isoenzymes. Further study showed that adrenaline significantly decreased serum albumin level and albumin-globulin ratio, while the level of APPs (α₁-acid glycoprotein and haptoglobin) is increased. The micronucleus test revealed a genotoxic effect of adrenaline at higher concentrations (1.5 mg/kg and 3 mg/kg body weight) compared to untreated rats. It can be concluded that adrenaline exerts oxidative and nitrative stress in rats, increased damage to lipids and proteins, and damage of cardiomyocytes and cytogenetic damage. Obtained results may contribute to better understanding of the toxicity of adrenaline with aims to preventing its harmful effects.

1. Introduction

Adrenaline (epinephrine) is a naturally occurring catecholamine which is secreted by the medulla of the adrenal glands. As a drug, adrenaline was discovered over a century ago and has been used in human cardiopulmonary resuscitation since 1922 [1]. Adrenaline also finds application in treatment of cardiac arrest, asthma, allergic reactions, and glaucoma [2]. Adrenaline acts through the alpha and beta adrenergic receptors which leads to vasoconstriction, an increase in the rate and force of contraction of the heart, and dilatation of bronchi and cerebral vessels. During normal physiological conditions, there is no constant secretion of adrenaline, but under the stress condition, a high level of adrenaline is released to prepare body for “fight or flight” response [3].

However, adrenaline at other catecholamine at doses exceeding physiological levels may cause toxic effects [4, 5]. There are studies indicating toxic effects of adrenaline via signal transduction pathways [6, 7]. Also, it seems that adrenaline exerts detrimental effects via oxidative products of its metabolism [8–10]. In line with this, it was shown that auto-oxidation of catecholamine may generate reactive oxygen...
species (ROS) [11]. One of the final products of oxidative metabolism—adrenochrome is able to stimulate oxidation of adrenaline in which superoxide anion ($O_2^-$) and hydrogen peroxide ($H_2O_2$) are formed. Besides, it was reported that catalase and quercetin diminished the DNA-damaging effect of adrenaline in vitro [12].

It is well known that overproduction of ROS can lead to disruption of redox balance and cause oxidative stress [13]. Namely, if antioxidative mechanisms fail to remove excess level of ROS, cells become prone to damage of DNA, proteins, and lipids [14]. Proteins are among major targets of ROS or reactive nitrogen species (RNS) and lead to changes in protein content such as carbonyl group formation and nitrotyrosine (3-NT) generation [15]. There are assumptions that high level of adrenaline may cause protein damage via free radicals [16]. It is worth noting that oxidative damage to proteins plays an important role in loss of physiological functions, thus favoring development of various diseases [17–19]. On the other hand, lipid peroxidation by ROS may initiate the peroxidation of membrane lipids and cause cellular injuries. In addition, oxidative DNA damages play a role in the mutagenesis and an increased risk of tumors [20, 21].

Although there are indications that adrenaline could increase the level of ROS, the cause–consequence relationship between them is not fully understood. For this purpose, we determined parameters of oxidative status: catalase (CAT), Cu, Zn-superoxide dismutase (SOD), malondialdehyde (MDA), nitrite ($NO_2^-$), lactate dehydrogenase (LDH), and carbonyl groups (PCC) after administration of adrenaline in Wistar rats. Also, we evaluated how adrenaline influenced the level of acute phase proteins (APPs), $\alpha_1$-acid glycoprotein (AGP), haptoglobin (Hp), and level of 3-NT. Possible genotoxic effects of adrenaline on bone marrow cells using micronucleus test were also estimated. The results of this investigation should contribute to a better understanding of adrenaline toxicity with aims to preventing its harmful effects.

2. Materials and Methods

2.1. Animals. In this study, Wistar rats aged 18 weeks weighing 220–280 g were obtained from the Institute of Biomedical Research, University of Belgrade. Three experimental groups with adrenaline and two control groups (negative and positive) contained seven male Wistar rats. The animals were kept under controlled constant environmental conditions (25 ± 4°C, 55 ± 5% humidity) with a 12/12 h light/dark cycle. They received food and water ad libitum. Care and animal treatment were conducted according to the Guide for the Care and Use of Laboratory Animals (National Research Council [22]). The investigation was approved by the Ethical Committee of the Faculty of Veterinary Medicine (University of Belgrade).

2.2. Doses and Treatment. Three experimental doses of adrenaline were used: 0.75 mg/kg, 1.5 mg/kg, and 3 mg/kg body weight, corresponding to 15%, 30%, and 60%, respectively, of LD50 [23]. Cyclophosphamide was used as a positive control and 60 mg/kg body weight per rats was given, while the negative control group was treated with saline (0.9% NaCl) [24]. The intraperitoneal (i.p.) route of application was used in all experimental groups. After 24 hours of treatments, all rats were euthanized. From each animal, blood and bone marrow from both femurs were taken.

2.3. Blood Sampling. Blood samples for the biochemical analysis were taken from the rats by puncture of v. jugularis and collected into heparinized tubes. Plasma was obtained from blood by centrifugation at 3000 rpm for 10 min. Erythrocytes were rinsed three times in isotonic solution of NaCl (0.9%). Then, samples were frozen at −20°C until analysis. Haemoglobin concentration was estimated by the cyanomethaemoglobin method [25]. Haemolysates of erythrocytes were used for determination of activities of antioxidant enzymes (SOD, CAT) and level of MDA. Plasma was used for determination of $NO_2^-$ concentration, 3-NT, protein carbonyl groups, total LDH level and its isoenzyme activity, and levels of APPs, Hp, and AGP.

2.4. Oxidative Stress Parameters. The Cu, Zn-superoxide dismutase (SOD) activity in erythrocytes was determined spectrophotometrically according to Misra and Fridovich [26] and expressed in units/gram of haemoglobin (U/g Hb). The relative activity of SOD was determined by means of vertical electrophoresis at 10% polyacrylamide gel electrophoresis (PAGE) and oxidation of nitro blue tetrazolium (NBT) following the Beauchamp and Fridovich method [27] (Hoeffer miniVE, Amersham, LKB, 2117, Bromma, Uppsala, Sweden). The band intensity was measured using TotalLab TL120, and results were expressed in arbitrary U/g Hb.

Catalase (CAT) activity in erythrocytes was assayed by means of the UV kinetic method at absorbance of 240 nm with the presence of $H_2O_2$ [28]. Activity was expressed as U/g Hb, calculated by using an extinction coefficient of 99.4 M$^{-1}$ cm$^{-1}$.

The level of malondialdehyde (MDA) in erythrocytes was quantified by measuring the formation of thiobarbituric acid reactive substances (TBARS) [29, 30]. The absorbance of the colored MDA-TBARS complex was measured at a wavelength of 535 nm and expressed in nM MDA/g Hb.

Nitrite concentration ($NO_2^-$) in plasma was determined with Griess reagent [31] on a microplate reader at 540 nm (plate reader, Mod. A1, Nubenco Enterprises, ICN). The results were expressed in µM.

The determination of carbonyl groups (PCC) was performed spectrophotometrically with 2,4-dinitrophenylhydroxazidine at 365 nm [32]. The concentration of carbonyl groups was calculated on the basis of the absorption coefficient for this chromogen ($a = 22,000$ M$^{-1}$ cm$^{-1}$; the results are shown in µM.

Alpha 1-acid glycoprotein (AGP) determined using Native PAGE, haptoglobin (Hp), and 3-nitrotyrosine (3-NT) levels in plasma of rats was determined using SDS-PAGE and Immunoblot methods with polyclonal antibody produced in rabbit (Sigma-Aldrich, St. Louise, USA) and monoclonal anti-3-nitrotyrosine antibody produced in mouse (Sigma-Aldrich, St. Louise, USA) [33, 34].
2.5. Determination of LDH. Total lactate dehydrogenase (LDH) was determined by following the initial rate of pyruvate reduction to lactate [34]. The activity of LDH was expressed in units per liter (U/L). Isoenzyme forms of LDH (LDH₁-LDH₅) were detected by PAGE technique (Hoeffer MiniVE, LKB, 2117, Bromma, Uppsala Sweden) using Tris-glycine buffer and sodium-lactate as substrates in the presence of nitro blue tetrazolium chloride [35]. Band intensity was measured densitometrically using TotalLab TL 120. The relative activity of isoenzymes was shown in percentages.

2.6. Native Gel Electrophoresis of Plasma Proteins. Vertical polyacrylamide gel electrophoresis at pH 8.6 (alkaline-PAGE) was carried out with Hoeffer miniVE cell (Amer sham, LKB, 2117, Bromma, Uppsala Sweden) at 120 V and room temperature for 2 h. The gel used (0.75 mm thick) consisted of 4.5% T staking gel and 8% T separation gel (T% is an expression representing the concentration of acrylamide plus bisacrylamide in the gel). The electrode and migration buffers consisted of 0.19 M glycine and 0.024 M Tris at pH 8.6. After electrophoresis, proteins were stained using Coomassie blue 0.1% [36]. The band intensity was measured using TotalLab TL 120. Results were shown in percentages in relation to the total area. The albumin:globulin ratio was calculated by dividing albumin content by the sum of α₁, β₁, and γ-globulins.

2.7. Micronucleus Test. The preparation and staining of bone marrow cells for the micronucleus test were performed according to Schmid [37] and Mavournin et al. [38]. After 24 hours of treatment, the bone marrow cells were flushed out with fetal calf serum, and the cells were suspended through centrifugation, smeared, and stained with May-Grünwald and Giemsa solution. A total of 1000 polychromatized erythrocytes were scored for each animal at a magnification of 100x (oil immersion) using a microscope (Leica, Germany). The PCE/NCE ratio was calculated to determine the cytotoxic effects of the adrenaline. All slides were coded and scored blind.

2.8. Statistical Analysis. Statistical significance of differences of all examined parameters was determined by means of the ANOVA test followed by Dunnett test. Normality tests were first performed for all groups using the d’Agostino-Pearson omnibus test. Data were expressed as means ± standard error (S.E.). Significance level was set at P < 0.05. Statistical analysis was done using GraphPad Prism 7.0 Software, CA, USA.

3. Results

Results of the analysis of total activity of Cu, Zn-SOD enzyme in rats treated with adrenaline are shown in Figure 1. It was observed that adrenaline treatment significantly decreased (P < 0.001) the total enzyme activity by 26.09%, 38.09%, and 69.97%, respectively, in relation to the control group. Cyclophosphamide, as positive control, also reduces activity of SOD (48.54%, P < 0.001). The decreased activities of SOD biochemical assay were verified by the results of electrophoresis as shown in Figure 1(b).

Significant increase in CAT activity was identified at all concentrations of adrenaline (Figure 2). The lowest concentration of adrenaline (0.75 mg/kg) induced significant increases (67.66 ± 5.83 U/g Hb, P < 0.05) in activity of CAT while the higher concentrations (1.5 mg/kg and 3 mg/kg) of adrenaline induce a more evident increase (92.50 ± 9.35 U/g Hb and 95.65 ± 9.73 U/g Hb, P < 0.001) compared to the control group (28.92 ± 1.83 U/g Hb). Similarly, the activity of CAT was significantly increased (P < 0.01) after treatment with cyclophosphamide.
In Figure 3, MDA levels after treatment with adrenaline are shown. Adrenalin exerted a significant increase in a dose-dependent manner (1.84-, 3.41-, 7.78-, and 5.29-fold, \( P < 0.001 \)) in MDA levels in rats compared to the untreated group (Figure 3). Also, cyclophosphamide induced a significant rise (\( P < 0.001 \)) in respect to the control.

Results of the total LDH and its relative isoenzyme distribution in rats treated with adrenaline are shown in Figures 4 and 5. There was clearly an increase (\( P < 0.001 \)) in total activity of LDH in the group treated with all doses of adrenaline compared to the control group. Cyclophosphamide induces a less profound effect than the group treated with 3.5 mg/kg dose of adrenaline, but the total activity of LDH is also significantly increased (\( P < 0.001 \)) in comparison to the control.

In Figure 5, it was observed that the LDH\(_1\) isoenzyme shows an evident increase (\( P < 0.001 \)) in rats treated with adrenaline in respect to the control group. The lower but statistically significant effect (\( P < 0.05, P < 0.01 \)) was evident in the activity of LDH\(_2\) isoenzyme after adrenaline treatment compared to the control group. Similarly, significant intensity bands for LDH\(_1\) and LDH\(_2\) isoenzymes were noticed in the group of rats treated with higher concentrations of adrenaline. The activity of isoenzymes LDH\(_1\) and LDH\(_2\) was also significantly increased (\( P < 0.01, P < 0.001 \)) in rats treated with cyclophosphamide.

The effect of adrenaline on PCC levels in rats is presented in Figure 6. Compared to untreated rats, the significant elevation (\( P < 0.001 \)) of PCC levels after treatment with all tested doses of adrenaline was noticed (Figure 6). Rats treated with
cyclophosphamide also showed a significant rise \((P < 0.001)\) in PCC levels in comparison to control.

The level of \(\text{NO}_2^-\) in rats after treatment with adrenaline is shown in Figure 7. Adrenaline induced a significant elevation of \(\text{NO}_2^-\) level in a dose-dependent manner. The highest concentration (3 mg/kg) of adrenaline induced the most distinct increases of \(\text{NO}_2^-\) \((P < 0.001)\) compared to the control group. Positive control also induces a significant increment in the level of \(\text{NO}_2^-\) \((P < 0.01)\).

The results of the electrophoretic distribution of plasma proteins are shown in Figure 8, while the results of their relative distribution are given in Table 1.

In our studies, reduced albumin concentration in adrenaline treatment has been demonstrated, and this decrease is dose-dependent. Adrenaline has shown to reduce albumin concentration by 5.81%, 16.09%, and 18.75% depending on the dose level, while due to exposure to cyclophosphamide, the decrease of albumin is 9.62%. By comparing the results of albumin level in all groups with the control group, the frequency of micronucleated polychromatized erythrocytes (MNPE) at higher concentrations (1.5 mg/kg and 3 mg/kg) of adrenaline decreases by 80.40% \((P < 0.05)\), 150.67% \((P < 0.01)\), and 79.36% \((P < 0.05)\). A significant decrease \((P < 0.05)\) of the \(\beta_1\)-fraction (transferrin, haemopexin, and \(\beta\)-lipoproteins) versus the control group was noticed in groups treated with adrenaline in doses of 0.75 mg/kg and 1.5 mg/kg \((P < 0.01)\), and 3 mg/kg \((P < 0.001)\), with increases by 42.97%, 45.09%, and 252.79%, respectively. The \(\alpha_2\)-fraction (Hp, ceruloplasmin, and \(\alpha_2\)-macroglobulin) was affected only by adrenaline treatment in doses of 0.75 mg/kg, 1.5 mg/kg, and 3 mg/kg and increases by 80.40% \((P < 0.05)\), 150.67% \((P < 0.01)\), and 79.36% \((P < 0.05)\). A significant decrease \((P < 0.05)\) of the \(\beta_1\)-fraction (transferrin, haemopexin, and \(\beta\)-lipoproteins) versus the control group was noticed in groups treated with adrenaline in doses of 0.75 mg/kg (18.88%), 1.5 mg/kg (27.63%), and 3 mg/kg (19.32%). The \(\beta_2\)-fraction (fibrinogen, C3, and \(\beta_2\)-microglobulin) in groups treated with adrenaline was decreased by 27.78% (dose 0.75 mg/kg), 28.24% (1.5 mg/kg), and 50.69% (3 mg/kg), which was significantly lower \((P < 0.001)\) than in the control group. In prealbumin and \(\gamma\)-globulin fractions, there were no significant differences between treated and control groups. A significant decrease \((P < 0.05)\) in the ratio \((A:G)\) versus control group was noticed in groups treated with adrenaline in doses of 1.5 mg/kg and 3 mg/kg.

The significant increase \((P < 0.001)\) in AGP levels was noticed in rats treated with adrenaline, especially at higher doses (1.5 mg/kg and 3 mg/kg) in respect to the control group (Figure 9). Related results of AGP levels on electrophoretogram were also detected.

The effect of adrenaline treatment on level of haptoglobin (Hp) in rats is presented in Figure 10. The significant difference \((P < 0.01, P < 0.001)\) was observed in the level of Hp in rats treated with 0.75 mg/kg, 1.5 mg/kg, and 3 mg/kg of adrenaline, versus the control group, respectively. These results conformed with the electrophoretic profile.

The effect of adrenaline treatment on level of 3-nitrotyrosine (3-NT) in rats is presented in Figure 11. A significant difference \((P < 0.001)\) was evident in the level of nitrotyrosine in rats treated with all doses (1.5 mg/kg and 3 mg/kg) of adrenaline while a 0.75 mg/kg dose of adrenaline affects \((P < 0.01)\) on the rise at the 3-NT level. In that manner, the intensity of the band was most evident at 3 mg/kg of adrenaline in the electrophoretic profile.

The results of the micronucleus (MN) test in bone marrow of rats treated with adrenaline are shown in Table 2. Adrenaline induced a significant increase \((P < 0.001)\) in the frequency of micronucleated polychromatized erythrocytes (MNPE) at higher concentrations (1.5 mg/kg, 3 mg/kg) when compared with the negative control group. Also, significant and dose-dependent decreases in polychromatized erythrocyte/normochromatic erythrocyte (PCE/NCE) ratio were seen in higher doses of adrenaline (1.5 mg/kg, 3 mg/kg). As expected, cyclophosphamide (60 mg/kg) induced a significant increase \((P < 0.001)\) in MNPE and decreases in the PCE/NCE ratio in respect to the control.

4. Discussion

In the last few decades, researchers focus on detecting harmful effects of chemical substances that are used as drugs in order to
activity in treated rats indicated that excess level of adrenaline may stimulate oxidation of adrenaline and thus Superoxide anions generated in oxidative metabolism of saturated fatty acids of the membrane are one of favored oxidation targets of ROS due to its oxygen-rich environment. In this study, SOD activity in erythrocytes was significantly increased in rats treated with adrenaline compared to the untreated group. SOD and CAT are the most important antioxidant enzymes in the defense system against ROS [39]. These results may indicate that administration of adrenaline in rats caused disruption in oxidant/antioxidant balance. Superoxide anions generated in oxidative metabolism of adrenaline may stimulate oxidation of adrenaline and thus increase the amount of free radicals [11]. Decreased SOD activity in treated rats indicated that excess level of superoxide anions induced attenuation of SOD activity. On the other hand, increased CAT activity found in rats treated with adrenaline implies that CAT has an important protective role in removing free radicals produced in cells. This assumption is supported by studies of Dželić et al. [10] and Radakovic et al. [12] where catalase in cotreatment with adrenaline reduced DNA damage mediated by free radicals -ect the activities of antioxidant enzymes.

In this study, SOD activity in erythrocytes was significantly decreased while CAT activity was significantly increased in rats treated with adrenaline compared to the untreated group. SOD and CAT are the most important antioxidant enzymes in the defense system against ROS [39]. These results may indicate that administration of adrenaline in rats caused disruption in oxidant/antioxidant balance. Superoxide anions generated in oxidative metabolism of adrenaline may stimulate oxidation of adrenaline and thus increase the amount of free radicals [11]. Decreased SOD activity in treated rats indicated that excess level of

| Groups | Prealbumin | Albumin | α1- | α2- | β1- | β2- | γ-Globulins | Ratio (A : G) |
|--------|------------|---------|-----|-----|-----|-----|-------------|--------------|
| C      | 2.66 ± 0.52 | 57.62 ± 3.80 | 5.19 ± 0.56 | 2.96 ± 1.62 | 15.89 ± 1.49 | 8.64 ± 0.79 | 7.04 ± 0.52 | 1.36 ± 0.22 |
| A1     | 4.57 ± 0.57** | 54.32 ± 1.07 | 7.42 ± 1.03** | 5.34 ± 1.20* | 12.89 ± 0.70* | 6.24 ± 1.27* | 9.78 ± 1.38* | 1.20 ± 0.15 |
| A2     | 3.66 ± 0.25 | 48.39 ± 3.72** | 8.53 ± 0.66** | 7.42 ± 1.36** | 11.50 ± 0.85** | 6.20 ± 1.28* | 9.43 ± 1.23 | 0.94 ± 0.22* |
| A3     | 2.05 ± 0.32 | 46.86 ± 2.54*** | 18.31 ± 3.01*** | 5.31 ± 1.32* | 12.82 ± 1.52* | 4.26 ± 2.57*** | 9.47 ± 1.56 | 0.88 ± 0.28* |
| CP     | 3.69 ± 0.47* | 52.07 ± 1.25 | 6.07 ± 1.73 | 2.17 ± 1.52 | 16.14 ± 0.83 | 9.66 ± 0.84** | 9.11 ± 1.42 | 1.08 ± 0.23 |

*P < 0.05, **P < 0.01, and ***P < 0.001. C: control group, groups treated with adrenaline doses (A1, 0.75 mg/kg; A2, 1.5 mg/kg; and A3, 3 mg/kg body weight); CP: group treated with cyclophosphamide.

Figure 9: Immunohistochemical detection of APG on Native PAGE. The level of AGP (a) with electrophoregram (b) after treatment with adrenaline. Control group - C, groups treated with adrenaline doses (A1-0.75 mg/kg; A2-1.5 mg/kg; A3-3 mg/kg body weight); group treated with cyclophosphamide (CP). Data are expressed as means ± SE. **P < 0.01, ***P < 0.001 vs. control group.

Figure 10: The level of haptoglobin (a) with electrophoregram (b) after treatment with adrenaline. Control group - C, groups treated with adrenaline doses (A1-0.75 mg/kg; A2-1.5 mg/kg; A3-3 mg/kg); group treated with cyclophosphamide (CP). Data are expressed as means ± SE. **P < 0.01, ***P < 0.001 vs. control group.
treated with inhibitors of free radicals. In addition, increased level of adrenaline. A similar effect that the detrimental effect was abolished when cells were treated with inhibitors of free radicals. In addition, increased ROS/RNS and increases protein oxidative damage considered line. They indicate that adrenaline stimulates production of ROS/RNS and increases protein oxidative damage consideration. 

Souza et al. [16] reported increased levels of carbonylated proteins, in plasma of rats treated with adrenaline. As a consequence of oxidative stress in rats treated with adrenaline. Namely, ONOO\(^-\) may spontaneously decompose to yield NO\(^2-\) and high reactive radical \("\)OH. Forming ONOO\(^-\) can take part of lipid peroxidation in reaction with unsaturated fatty acid-containing liposomes. In addition, ONOO\(^-\) may influence on protein participation in signal transduction mechanisms [48]. It was found that high concentrations of ONOO\(^-\) often lead to necrotic-type cell death [49]. Supportive evidence about the presence of nitrosative stress is revealed by our findings that an increase in the level of 3-NT was detected in the adrenaline-treated group. Consistent with our result is the study of Romana-Souza et al. [16] who reported an increased level of nitrotyrosine in mouse skin after treatment with a high level of adrenaline. The mechanism of protein tyrosine nitration in biological systems has been well described [50]. Nitrotyrosine formation has been observed in various cardiovascular diseases [50–52]. An increased level of 3-NT may cause alternation of protein function, protein-protein interactions, and cell signaling [50, 53, 54]. Also, one of the consequences of adrenaline application is the acute phase response, which we demonstrated by evaluating APPs. One of the most important metabolic changes during the acute phase is the production of APPs which are released from the liver into the plasma [55]. This occurs within a few hours, and these proteins remain elevated as long as the inflammatory stimulus persists, making them perfect indicators of inflammation or injury, and useful for predicting the outcome of disease. Their only flaw is that they are poorly specific. APPs have been widely used in human and veterinary medicine as biomarkers of diseases, inflammatory processes, and various infections [56, 57]. Classification of APPs can be done according to the magnitude of the increase (positive APPs) or decrease (negative APPs) in their serum concentrations during the acute phase response [58]. Some of the APPs (\(\alpha_1\)-antitrypsin, and \(\alpha_2\)-macroglobulin) have antiprotease activity designed to inhibit phagocyte proteases and to minimize tissue damage. Others (\(\alpha_1\)-acid glycoprotein) have antibacterial or scavenging activity (haptoglobin, serum amyloid A, and C-reactive protein), by binding metabolites released from cellular degradation so they cannot be utilized by pathogen. Albumin, as a negative APP, is a major source of amino acids and is responsible for about 75% of the osmotic pressure of plasma. In this study, it has been shown that adrenaline reduced the albumin concentration by a dose-dependent level and also exposure to cyclophosphamide led to a decrease of albumin. Adrenaline causes an increased flux of free radicals, which can be due to the oxidation of thiol groups and the formation of albumin dimers or polymers, and consequently the reduction in osmotic pressure a c.

![Figure 11: The level of nitrotyrosine (a) with electrophoregram (b) after treatment with adrenaline. Control group - C, groups treated with adrenaline doses (A1-0.75 mg/kg; A2-1.5 mg/kg; A3-3 mg/kg body weight); group treated with cyclophosphamide (CP). Data are expressed as means ± SE. **P < 0.01, ***P < 0.001 vs. control group.](image-url)
The effect of the acute phase protein AGP in rats exposed to adrenaline was also estimated. A significant dose-dependent elevation in AGP following adrenalin treatment was observed. AGP is a positive APP with a normal concentration in the human plasma at 0.6–1.2 mg/mL [59]. The plasma concentration of AGP can increase from 2- to 10-fold when influenced by various factors, such as stress, inflammation drugs (phenobarbitone and rifampicin), burns, infections, and pregnancy. AGP possesses immunomodulatory activities and modulates neutrophil chemotactic migration and superoxide generation [60]. AGP significantly suppresses synthesis of IL-2, proliferation of lymphocytes, and platelet aggregation [61]. It has been shown that AGP protects neutrophil generation of ROS probably due to binding of free radicals [62]. AGP can bind to a number of metabolites such as heparin, histamine and serotonin, steroids, catecholamines, and pharmaceutical compounds. Increased AGP may affect pharmacokinetics by reducing the concentration of free drugs, by binding to the metabolically active fraction of the drug. Matsumoto et al. [63] reported that human AGP at physiological concentrations protects erythrocytes from \( \text{H}_2\text{O}_2 \). On the basis of the above-mentioned, we assume that the AGP level increases in response to oxidative stress after treatment with adrenaline. In this study, we detected an increased Hp level in rats treated with higher doses of adrenaline (1.5 and 3 mg/kg). Haptoglobin (Hp) is a positive acute-phase glycoprotein classified in α2 fraction together with fibrinogen and α-globulins with antiprotease activity [64]. Hp has a pronounced anti-inflammatory action, because of its ability to bind to heme of haemoglobin, forming an Hp-Hb complex. By forming this complex, Hp prevents the promotion of free radicals and its accumulation in endothelial cells which is catalysed by heme. Free Hb has the ability to catalyse the formation of hydroxyl radicals from \( \text{H}_2\text{O}_2 \). By binding to neutrophils, Hp may inhibit NADPH oxidase activation and therefore the production of reactive forms of oxygen associated with inflammation. The Hp-Hb complex, by removing free Hb, prevents renal injury that may occur when free Hb passes through the glomerular filter [65]. There is a great variability in antioxidant capacity, depending on Hp polymorphism, so individuals with Hp2-2 have a lower antioxidant capacity [65]. The Hp-Hb complex also binds nitric oxide or nitrogen monoxide (NO) [66]. This action is also phenotype-dependent. Due to its longer half-life, the Hp2-2/Hb complex scavenges more NO than the Hp1-1/Hb complex does [67, 68]. Haptoglobin can also modulate the immune response by binding to receptors on immune cells. Glucocorticoids and catecholamines activate haptoglobin synthesis whereas insulin exerts an opposite action. Since increased levels of extracellular Hb can occur due to impaired membrane integrity, we assumed that Hp scavenges free Hb as a result of lipid peroxidation. This fact coincides with our results of MDA analysis and leads to a conclusion that an elevated Hp level has a protective response to adrenaline-induced oxidative stress. Our findings unequivocally point that adrenaline induces an acute phase response. Although the APPs are a significant marker of inflammation and/or infection, it seems that these results give a new linkage between APPs and noninflammatory stress.

The possible genotoxicity of adrenaline was evaluated by a micronucleus test in bone marrow cells of rats. The results indicate that adrenaline expresses a genotoxic effect at higher concentrations (1.5 and 3 mg/kg) since it caused a significant induction of MN in the bone marrow of rats. This result is in compliance with the study of McGregor et al. [69] in which the adrenaline exhibited a mutagenic effect on mouse L5178Y lymphoma cells. In the Comet assay, adrenaline induced DNA damage in 3T3 cells of rats [6]. In our work, tested catecholamine (1.5 and 3 mg/kg) decreased the PCE/NCE ratio indicating its cytotoxicity in bone marrow. Muthuraman et al. [70] showed that adrenaline possessed a cytotoxic effect and affects DNA fragmentation in a dose-dependent manner in MDCK cells.

Several studies have implicated involvement of free radicals in the genotoxic action of adrenaline [12, 71]. Antioxidant enzymes CAT and SOD are the integral part of antioxidant defense mechanisms and have a considerable importance since they are involved in protection from free radicals. We assume that the antioxidants respond to oxidative stress caused by adrenaline since significant changes in the CAT and SOD activity in treated rats were detected. Martinez et al. [72] have classified catecholamines as potent oxidative mutagens. It has been suggested that catecholamine generates free radicals by autoxidation and redox cycling [44]. ROS have the ability to induce various types of DNA damage such as double-strand breaks (DSB), and this type of DNA damage is considered as a main contributor to MN induction [73]. We suggest that the increased induction of MN in the bone marrow of rats is a result of increased production of free radicals produced by oxidative metabolism of adrenaline. So, it should be expected that antioxidants could protect cells from toxic effects of adrenaline.

| Treatment          | Treatment time (h) | Doses mg/kg | Total cell number | MNPCE (%) ± SE | PCE (%) ± SE |
|--------------------|--------------------|-------------|-------------------|----------------|--------------|
| Negative control   | 24                 | 0           | 5000              | 0.80 ± 0.37    | 49.24 ± 0.02 |
| Adrenaline         | 24                 | 0.75        | 5000              | 1.20 ± 0.49    | 47.92 ± 0.21 |
|                    | 24                 | 1.5         | 5000              | 2.80 ± 0.37*   | 46.40 ± 0.44** |
|                    | 24                 | 3           | 5000              | 5.40 ± 0.24*   | 43.43 ± 1.19*** |
| Cyclophosphamide   | 24                 | 60          | 5000              | 14.40 ± 0.40*  | 47.94 ± 0.38*** |

Data are expressed as means ± SE. **P < 0.01 and ***P < 0.001 vs. the control group.
5. Conclusion

In summary, our results show that adrenaline induces oxidative and nitritative stress in Wistar rats, accompanied by changes in the activity of antioxidant enzymes, increased damage to lipids and proteins, increased level of NO2 and nitrotyrosine derivate, damage of cardiomyocytes, and genotoxic damage. Also, adrenaline exerts acute-phase response through increased level of AGP and Hp and decreased level of serum albumin level. Therefore, obtained results may contribute to better understanding of adrenaline toxicity with aims at preventing its harmful effects.

Data Availability

The data used to support the findings of this study are included within the article.

Conflicts of Interest

All authors declare that there are no conflicts of interest.

Acknowledgments

This work was funded by the Ministry of Education, Science and Technological Development of Serbia (Grants: OI173034 and III46002).

References

[1] J. A. Cooper, J. D. Cooper, and J. M. Cooper, “Cardiopulmonary resuscitation: history, current practice, and future direction,” Circulation, vol. 114, no. 25, pp. 2839–2849, 2006.

[2] D. Dietz, Toxicology and Carcinogenesis Studies of Epinephrine Hydrochloride (CAS NO. 59-31-2) in F344/N Rats and B6C3F1 Mice (Inhalation Studies), National Toxicology Program, Research Triangle Park, NC, USA, 1990.

[3] L. Gavrilovic, V. Stojiljkovic, J. Kasapovic et al., “Chronic physical stress changes gene expression of catecholamine biosynthetic enzymes in the adrenal medulla of adult rats,” Acta Veterinaria-Beograd, vol. 62, no. 2-3, pp. 151–169, 2012.

[4] G. S. Behonick, M. J. Novak, E. W. Nealley, and S. I. Baskin, “Toxicology update: the cardiotoxicity of the oxidative stress metabolites of catecholamines (aminochromes),” Journal of Applied Toxicology, vol. 21, no. S1, pp. S15–S22, 2001.

[5] V. M. Costa, R. Silva, L. M. Ferreira et al., “Oxidation process of adrenaline in freshly isolated rat cardiomyocytes: formation of adenochrome, quinoproteins, and GSH adduct,” Chemical Research in Toxicology, vol. 20, no. 8, pp. 1183–1191, 2007.

[6] M. S. Flint, A. Baum, W. H. Chambers, and F. I. Jenkins, “Induction of DNA damage, alteration of DNA repair and transcriptional activation by stress hormones,” Psychoneuroendocrinology, vol. 32, no. 5, pp. 470–479, 2007.

[7] A. Adamoeva, Y. Abdellatif, and N. S. Dhalla, “Role of the excessive amounts of circulating catecholamines and glucocorticoids in stress-induced heart disease,” Canadian Journal of Physiology and Pharmacology, vol. 87, no. 7, pp. 493–514, 2009.

[8] T. Miura, S. Muraoka, Y. Fujimoto, and K. Zhao, “DNA damage induced by catechol derivatives,” Chemico-Biological Interactions, vol. 126, no. 2, pp. 125–136, 2000.

[9] N. S. Dhalla, H. Sasaki, S. Mochizuki, K. S. Dhalla, X. Liu, and V. Elminab, “Catecholamine-induced cardiomyopathy,” in Cardiovascular Toxicity, D. Acosta, Ed., pp. 269–318, Raven Press, New York, NY, USA, 2001.

[10] N. Djelic, M. Radakovic, B. Spremo-Popavec et al., “Evaluation of cytogenetic and DNA damage in human lymphocytes treated with adrenaline in vitro,” Toxicology In Vivo, vol. 29, no. 1, pp. 27–33, 2015.

[11] M. L. Genova, N. M. Abd-Elsalam, E. S. M. E. Mahdy et al., “Redox cycling of adrenaline and adrenochrome catalysed by mitochondrial complex I,” Archives of Biochemistry and Biophysics, vol. 447, no. 2, pp. 167–173, 2006.

[12] M. Radakovic, N. Djelic, J. Stevanovic et al., “The investigation of DNA damage induced by adrenaline in human lymphocytes in vitro/Ispitivanja Ostećenja DNK Ispazavanih Adrenalinom U Limfocitima Coveka u vitro,” Acta Veterinaria, vol. 64, no. 3, pp. 281–292, 2014.

[13] B. Halliwell and J. M. C. Gutteridge, “Role of free radicals and catalytic metal ions in human disease: an overview,” Methods in Enzymology, vol. 186, pp. 1–85, 1990.

[14] A. Rahal, A. Kumar, V. Singh et al., “Oxidative stress, prooxidants, and antioxidants: the interplay,” BioMed Research International, vol. 2014, Article ID 761264, 19 pages, 2014.

[15] E. Miller, A. Walczak, J. Saluk, M. B. Ponczek, and I. Majsterek, “Oxidative modification of patient’s plasma proteins and its role in pathogenesis of multiple sclerosis,” Clinical Biochemistry, vol. 45, no. 1–2, pp. 26–30, 2012.

[16] B. Romana-Souza, G. Santos Lima-Cezar, and A. Monte-Alto-Costa, “Psychological stress-induced catecholamines accelerates cutaneous aging in mice,” Mechanisms of Ageing and Development, vol. 152, pp. 63–73, 2015.

[17] R. T. Dean, R. Dunlop, P. Hume, and K. J. Rodgers, “Proteolytic ‘defences’ and the accumulation of oxidized polypeptides in cataractogenesis and atherogenesis,” Biochemical Society Symposium, vol. 70, pp. 135–146, 2003.

[18] E. R. Stadtman, “Protein oxidation and aging,” Free Radical Research, vol. 40, no. 12, pp. 1250–1258, 2006.

[19] Q. Ding, E. Dimayuga, and J. Keller, “Oxidative damage, protein synthesis, and protein degradation in Alzheimers disease,” Current Alzheimer Research, vol. 4, no. 1, pp. 73–79, 2007.

[20] S. Loff and H. E. Poulsen, “Cancer risk and oxidative DNA damage in man,” Journal of Molecular Medicine, vol. 74, no. 6, pp. 297–312, 1996.

[21] J. E. Klaunig and L. M. Kamendulis, “The role of oxidative stress in carcinogenesis,” Annual Review of Pharmacology and Toxicology, vol. 44, no. 1, pp. 239–267, 2004.

[22] National Research Council, Guide for the Care and Use of Laboratory Animals, National Academies Press, 2010.

[23] C. D. Barnes and L. G. Eltherington, “Carcinogenicity of synthetic enzymes in the adrenal medulla of adult rats,” Acta Veterinaria-Beograd, vol. 45, no. 1–2, pp. 23–28, 2004.

[24] D. Anderson, J. B. Bishop, R. C. Garner, P. Ostrosky-Wegman, and J. E. Klaunig, “Carcinogenicity of synthetic enzymes in the adrenal medulla of adult rats,” Acta Veterinaria-Beograd, vol. 45, no. 1–2, pp. 29–38, 2004.
[25] L. Tentori and A. M. Salvati, "Hemoglobinometry in human blood," *Methods in Enzymology*, vol. 76, pp. 707–715, 1981.

[26] H. P. Misra and I. Fridovich, "The role of superoxide anion in the autoxidation of epinephrine and a simple assay for superoxide dismutase," *The Journal of Biological Chemistry*, vol. 247, no. 10, pp. 3170–3175, 1972.

[27] C. Beauchamp and I. Fridovich, "Superoxide dismutase: improved assays and an assay applicable to acrylamide gels," *Analytical Biochemistry*, vol. 44, no. 1, pp. 276–287, 1971.

[28] H. Aebi, "Catalase in vitro," in *Methods in Enzymology*, L. Packer, Ed., pp. 121–126, Academic Press, Orlando, FL, USA, 1984.

[29] J. M. Gutteridge, "Lipid peroxidation and antioxidants as biomarkers of tissue damage," *Clinical Chemistry*, vol. 41, 12, Part 2, pp. 1819–1828, 1995.

[30] N. Traverso, S. Menini, E. P. Mainieri et al., "Malondialdehyde, a lipoperoxidation-derived aldehyde, can bring about secondary oxidative damage to proteins," *The Journals of Gerontology Series A: Biological Sciences and Medical Sciences*, vol. 59, no. 9, pp. B890–B895, 2004.

[31] I. Guevara, J. Iwanenko, A. Dembinska-Kiec et al., "Determination of nitrite/nitrate in human biological material by the simple Griess reaction," *Clinica Chimica Acta*, vol. 274, no. 2, pp. 177–188, 1998.

[32] R. L. Levine, D. Garland, C. N. Oliver et al., "Determination of carbonyl content in oxidatively modified proteins," *Methods in Enzymology*, vol. 186, pp. 464–478, 1990.

[33] H. Towbin, T. Staehelin, and J. Gordon, "Electrophoretic transfer of proteins from polyacrylamide gels to nitrocellulose sheets: procedure and some applications," *Proceedings of the National Academy of Sciences of the United States of America*, vol. 76, no. 9, pp. 4350–4354, 1979.

[34] T. Mahmood and P. C. Yang, "Western blot: technique, theory, and trouble shooting," *North American Journal of Medical Sciences*, vol. 4, no. 9, pp. 429–434, 2012.

[35] M. Yoshida and Y. Takakuwa, "Method for the simultaneous assay of initial velocities of lactate dehydrogenase isoenzymes following gel electrophoresis," *Journal of Biochemical and Biophysical Methods*, vol. 34, no. 3, pp. 167–175, 1997.

[36] B. D. Hames and D. Rickwood, *Gel Electrophoresis of Proteins, a Practical Approach*, Oxford University Press, New York, NY, USA, 1990.

[37] W. Schmid, "The micronucleus test," *Mutation Research/Environmental Mutagenesis and Related Subjects*, vol. 31, no. 1, pp. 9–15, 1975.

[38] K. H. Mavournin, D. H. Blakey, M. C. Cimino, M. F. Salamone, and J. A. Heddle, "The in vivo micronucleus assay in mammalian bone marrow and peripheral blood. A report of the U.S. Environmental Protection Agency Gene-Tox Program," *Mutation Research/Reviews in Genetic Toxicology*, vol. 239, no. 1, pp. 29–80, 1990.

[39] I. Fridovich, "Superoxide radical and superoxide dismutases," *Annual Review of Biochemistry*, vol. 64, no. 1, pp. 97–112, 1995.

[40] B. Pereira, L. F. B. P. Costa-Rosa, E. J. H. Bechara, P. Newsholme, and R. Curi, "Changes in the TBARS content and superoxide dismutase, catalase and glutathione peroxidase activities in the lymphoid organs and skeletal muscles of adrenomedullated rats," *Brazilian Journal of Medical and Biological Research*, vol. 31, no. 6, pp. 827–833, 1998.

[41] E. N. Frankel, "Chemistry of free radical and singlet oxidation of lipids," *Progress in Lipid Research*, vol. 23, no. 4, pp. 197–221, 1984.

[42] B. Bukowska, J. Michalowicz, and A. Marczak, "The effect of catechol on human peripheral blood mononuclear cells (in vitro study)," *Environmental Toxicology and Pharmacology*, vol. 39, no. 1, pp. 187–193, 2015.

[43] M. Preus, A. Bhargava, A. Elrahmankhater, and P. Gunzel, "Diagnostic value of serum creatine kinase and lactate dehydrogenase isoenzyme determinations for monitoring early cardiac damage in rats," *Toxicology Letters*, vol. 42, no. 2, pp. 225–233, 1988.

[44] A. Bindoli, D. J. Deeble, M. P. Rigobello, and L. Galzigna, "Direct and respiratory chain-mediated redox cycling of adrenochrome," *Biochimica et Biophysica Acta (BBA) - Bioenergetics*, vol. 1016, no. 3, pp. 349–356, 1990.

[45] B. Halliwell and M. Whiteman, "Measuring reactive species and oxidative damage in vivo and in cell culture: how should you do it and what do the results mean?", *British Journal of Pharmacology*, vol. 142, no. 2, pp. 231–255, 2004.

[46] J. Zweier and M. Talukder, "The role of oxidants and free radicals in reperfusion injury," *Cardiovascular Research*, vol. 70, no. 2, pp. 181–190, 2006.

[47] J. Stamler, D. Singel, and J. Loscalzo, "Biochemistry of nitric oxide and its redox-activated forms," *Science*, vol. 258, no. 5090, pp. 1898–1902, 1992.

[48] N. Campolo, S. Bartesaghi, and R. Radi, "Metal-catalyzed protein tyrosine nitration in biological systems," *Redox Report*, vol. 19, no. 6, pp. 221–231, 2014.

[49] L. Virág, E. Szabó, and P. Gergely, "C. Szabo "Peroxyxinitrite-induced cytotoxicity: mechanism and opportunities for intervention," *Toxicology Letters*, vol. 140–141, pp. 113–124, 2003.

[50] P. Pacher, J. S. Beckman, and L. Liaudet, "Nitric oxide and peroxynitrite in health and disease," *Physiological Reviews*, vol. 87, no. 1, pp. 315–424, 2007.

[51] C. Vadseth, J. M. Souza, L. Thomson et al., "Pro-thrombotic state induced by post-translational modification of fibrinogen by reactive nitrogen species," *Journal of Biological Chemistry*, vol. 279, no. 10, pp. 8820–8826, 2004.

[52] S. Pennathur, C. Bergt, B. Shao et al., "Human atherosclerotic intima and blood of patients with established coronary artery disease contain high density lipoprotein damaged by reactive nitrogen species," *Journal of Biological Chemistry*, vol. 279, no. 41, pp. 42977–42983, 2004.

[53] T. Koeck, X. Fu, S. L. Hazen, J. W. Crabb, D. J. Stuehr, and K. S. Aulak, "Rapid and selective oxygen-regulated protein tyrosine denitration and nitration in mitochondria," *Journal of Biological Chemistry*, vol. 279, no. 26, pp. 27257–27262, 2004.

[54] L. A. Abriata, A. Cassina, V. Tórtora et al., "Nitrination of solvent-exposed tyrosine 74 on cytochrome c triggers heme iron-methionine 80 bond disruption: nuclear magnetic resonance and optical spectroscopy studies," *Journal of Biological Chemistry*, vol. 284, no. 1, pp. 17–26, 2009.

[55] H. Baumann and J. Gauldie, "The acute phase response," *Immunology Today*, vol. 15, no. 2, pp. 74–80, 1994.

[56] P. M. Ridker, "Inflammatory biomarkers and risks of myocardial infarction, stroke, diabetes, and total mortality: implications for longevity," *Nutrition Reviews*, vol. 65, no. 12, pp. 253–259, 2007.
Oxidative Medicine and Cellular Longevity

[57] P. D. Eckersall and R. Bell, “Acute phase proteins: biomarkers of infection and inflammation in veterinary medicine,” *The Veterinary Journal*, vol. 185, no. 1, pp. 23–27, 2010.

[58] H. H. Petersen, J. P. Nielsen, and P. M. H. Heegaard, “Application of acute phase protein measurements in veterinary clinical chemistry,” *Veterinary Research*, vol. 35, no. 2, pp. 163–187, 2004.

[59] S. Colombo, T. Buclin, L. A. Décoyder et al., “Orosomucoid (α1-acid glycoprotein) plasma concentration and genetic variants: effects on human immunodeficiency virus protease inhibitor clearance and cellular accumulation,” *Clinical Pharmacology & Therapeutics*, vol. 80, no. 4, pp. 307–318, 2006.

[60] T. Hochepied, F. G. Berger, H. Baumann, and C. Libert, “α1-acid glycoprotein: an acute phase protein with inflammatory and immunomodulating properties,” *Cytochrome & Growth Factor Reviews*, vol. 14, no. 1, pp. 25–34, 2003.

[61] S. A. Elg, A. R. Mayer, L. F. Carson, L. B. Twiggs, R. B. Hill, and A. Suenaga, “OxyR, uvrA and its oxidative mutagens,” *Mutation Research/Genetic Toxicology and Environmental Mutagenesis*, vol. 467, no. 1, pp. 41–53, 2000.

[62] F. van Goethem, D. Lison, and M. Kirsch-Volders, “Comparative evaluation of the in vitro micronucleus test and the alkaline single cell gel electrophoresis assay for the detection of DNA damaging agents: genotoxic effects of cobalt powder, tungsten carbide and cobalt–tungsten carbide,” * Mutation Research/Genetic Toxicology and Environmental Mutagenesis*, vol. 392, no. 1-2, pp. 31–43, 1997.

[63] A. L. Pukhal’skii, G. V. Shmarina, E. A. Kalashnikova et al., “Effect of semisynthetic analog of α1-acid glycoprotein on immunomodulatory and antiinflammatory activity of natural glycoproteinglycoprotein on immunomodulatory and anti-inflammatory activity of natural glycoprotein,” *Bulletin of Experimental Biology and Medicine*, vol. 129, no. 5, pp. 480–483, 2000.

[64] K. Matsumoto, K. Nishi, Y. Tokutomi, T. Irie, A. Suegawa, and M. Otagiri, “Effects of α1-acid glycoprotein on erythrocyte deformability and membrane stabilization,” *Biological and Pharmaceutical Bulletin*, vol. 26, no. 1, pp. 123–126, 2003.

[65] E. Gruys, M. J. M. Toussaint, T. A. Niewold, and S. J. Koopmans, “Acute phase reaction and acute phase proteins,” *Journal of Zhejiang University Science*, vol. 6B, no. 11, pp. 1045–1056, 2005.

[66] S. Fagonee, J. Gburek, E. Hirsch et al., “Plasma protein haptoglobin modulates renal iron loading,” *The American Journal of Pathology*, vol. 166, no. 4, pp. 973–983, 2005.

[67] I. Azarov, X. He, A. Jeffers et al., “Rate of nitric oxide scavenging by hemoglobin bound to haptoglobin,” *Nitric Oxide*, vol. 18, no. 4, pp. 296–302, 2008.

[68] A. P. Levy, R. Asleh, S. Blum et al., “Haptoglobin: basic and clinical aspects,” *Antioxidants & Redox Signaling*, vol. 12, no. 2, pp. 293–304, 2010.

[69] A. I. Alayash, “Haptoglobin: old protein with new functions,” *Clinica Chimica Acta*, vol. 412, no. 7-8, pp. 493–498, 2011.

[70] D. B. McGregor, C. G. Riach, A. Brown et al., “Reactivity of catecholamines and related substances in the mouse lymphoma L5178Y cell assay for mutagens,” *Environmental and Molecular Mutagenesis*, vol. 11, no. 4, pp. 523–544, 1988.

[71] P. Muthuraman, P. C. Nagajyothis, M. Chandrasekaran et al., “Differential sensitivity of Madin-Darby canine kidney (MDCK) cells to epinephrine,” *The Journal of Nutrition, Health & Aging*, vol. 20, no. 5, pp. 486–493, 2016.

[72] M. E. Crespo and M. P. Bicho, “Membrane-mediated effects of catecholamines on the DNA of human leukocytes: the role of reactive oxygen species,” *Neurosignals*, vol. 4, no. 2, pp. 78–85, 1995.

[73] A. Martinez, A. Urios, and M. Blanco, “Mutagenicity of 80 chemicals in *Escherichia coli* tester strains IC203, deficient in OxyR, and its oxyR” parent WP2 uvrA/pKM101: detection of 31 oxidative mutagens,” *Mutation Research/Genetic Toxicology and Environmental Mutagenesis*, vol. 467, no. 1, pp. 41–53, 2000.