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Changes in oxidative markers in COVID-19 patients

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Aim and Background. Covid-19 has been as an important human infectious disease that has affected several countries. Cytokine storm has major role is Covid-19 pathogenesis. The association between inflammation and oxidative stress is well established. In this article, we aim to assess oxidative stress markers in Covid-19 patients compare to the healthy subjects.

Method. A total of 48 persons (24 with Covid-19 and 24 controls) were evaluated in this research. Serum oxidative stress markers including Malondialdehyde (MDA), total oxidant status (TOS), activity of catalase (CAT) and super oxide dismutase (SOD) were measured alongside routine laboratory tests.

Results. Patients group were divided into ICU and Non-ICU groups. ESR, CRP and serum level of ferritin were significantly higher in case group. Serum level of albumin was significantly lower in Covid-19 patients. Serum MDA and TOS was significantly increased in Covid-19 patients. Also, Covid-19 patients had higher serum activity of CAT and GPX.

Conclusion. Oxidative stress markers are significantly elevated in Covid-19 patients. This may have significant role in mechanism of disease development. In the fight against Covid-19, as a global struggle, all possible treatments demand more attention. So, Covid-19 patients may benefit from strategies for reducing or preventing oxidative stress.

Keywords: Covid-19, oxidative stress, inflammation, pathogenesis.

Introduction

Coronavirus disease 2019 (Covid-19) pandemic has rapidly spread around the world and encountered as a significant threat to global health. This disease identified for the first time in December 2019 in Wuhan, China (1). The severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2), an enveloped RNA betacoronavirus, is pathogenic agent of Covid-19 disease (2). Easily transmission of SARS-CoV-2 has major role in widely spread of this viral infection (3). The clinical course of Covid-19 may vary from asymptomatic to severe symptoms (acute lung injury), hospitalization for 3-4 d, transfer to intensive care unit (ICU) and even death (4). Upon the past months, Covid-19 crisis has dramatically increased risk of detrimental outcomes. Therefore, it is required to identify complicated pathogenic mechanisms of disease to decrease the risk of hospitalization and mortality. Commonly, respiratory viral infections have close association with inflammation and cytokine production (5). High serum level of cytokines and chemokines are reported in Covid-19 patients (6). In some cases, cy-
Cytokine storm developed which is regarded as a key factor in acute respiratory distress syndrome development and various organ dysfunctions (7). Oxidative stress has critical function in inflammatory processes; reactive oxygen species (ROS) and H2O2 can activate NF-κB to trigger inflammatory cytokine production (8). It is postulated that oxidative stress is associated with pathogenesis of SARS-CoV-2 infection (9). In this research, we measured oxidative markers including TOS (total oxidant status), MDA (Malondialdehyde) and antioxidant markers including CAT (catalase), SOD (super oxide dismutase) in covid-19 patients compared to healthy control subjects.

Method and Material

Patients and Informed Consent

This case-control study was conducted at the public hospitals of Hamadan province in Iran, which were designated for Covid-19 patients. Twenty-four covid-19 patients (female: male; 17:7) and 24 healthy subjects (female: male; 14:10) were enrolled. The inclusion criteria for the control and case group were as follows: healthy subjects without any underlying condition and patients with laboratory-confirmed Covid-19 without any underlying condition, respectively. The Covid-19 diagnosis was based on World Health Organization interim guidelines (10) infection is suspected: interim guidance 2020. Subjects with a history of diabetes, hypertension, cancers, auto immune disorders, and smokers were excluded both from control and case group. Also, subjects excluded if they had special regimen or take antioxidant supplements such as vitamin C, vitamin E, Coenzyme Q10, selenium, and so on. The protocol of this study was approved by Hamadan University of Medical Sciences with code 9906113797 and ethical code: IR.UMSHA.REC.1399.441. All patients gave their informed consent before being included in the investigation.

Sample Collection

Venous blood samples were collected in any time of the day within 24 h after admission. The samples were centrifuged at 3000g for 10 min; serum was separated, aliquot and stored at −20 C till analyzed.

Biochemical Assays

After blood samples collection, routine laboratory tests, such as CBC, C-reactive protein (CRP), ESR were done. Also, serum Ferritin level was determined by Pishgaman Sanjesh kit. Serum albumin was measured by Man company kit. CRP was measured by a semi-quantitative latex agglutination technique (Bionik slide agglutination test kit). ESR was quantified by a SE 9020 analyzer and CBC was determined by a Sysmex (Kolbe, Japan) SE 9020 analyzer.

Measurement of Oxidative Stress Markers

Serum MDA level measurement was based on Yagi method. Thiobarbituric acid (TBA) reactive substances formation, as a reaction product, was measured at a wavelength of 532 nm using spectrophotometry (11). The serum level of TOS was evaluated according to Erel, s method (12). SOD and CAT serum level were measured using the commercially available ELISA kits according to the instructions of the company (KIAZIST Life Sciences, Iran. Cat. No: KSOD96 and KCAT96, respectively).

Statistical Analysis

The distribution of qualitative data was described by frequencies and percentages and compared between three groups (control, Non-ICU, and ICU patients) by Fisher exact test. The quantitative data were described as the mean ± standard deviation (SD) and their Normal distribution was evaluated by Shapiro-Wilk test. In case of normal distribution, the mean of quantitative data was compared in three groups using one-way Analysis of Variance (ANOVA). Moreover, the pairwise comparisons were done using Tuckey post hoc test. All analyses were performed at 0.05 significance levels using SPSS version 23 (SPSS Inc., USA) and GraphPad Prism version 6 for Windows.

Result

In accordance with clinical features and intensive care unit (ICU) admission, case group were divided into two groups: ICU and Non-ICU patients. The mean ± SD age of the control group (51.5 ± 13.58 years), Non-ICU (57.36 ± 10.87 years) and ICU patients (55.5 ± 15.65 years.) was not statistically significant (p >0.05). The main characteristics of case (ICU and Non-ICU) and control groups are shown in Table 1. All subjects were comparable for age and sex. The control group was medication free, but all Covid-19 patient received Acetaminophen codeine, enoxaparin, azitromicina, dexamethasone, dextromethorphan, kalastra, naproxen, Vitamin D, and pantoprazole. Ten patients required ICU admission and treatment. The common symptoms on admission were fever (79.6%), cough (62.5%), respiratory distress (79%) and oxygen saturation (Spo2) less than 88 (75%).

Some of the biochemical and hematological characteristics are shown in Table 2. The mean level of ESR, CRP, Ferritin and Neutrophil count were significantly highest in ICU group and the lowest in control group (p <0.001). In addition, a significant difference was observed between the mean level of albumin and Lymphocyte count between
these three groups: the lowest level in ICU group and highest level in control group. Moreover, no significant differences were observed in WBC, hemoglobin, and platelet count between these groups. Pairwise comparison of significant characteristics is indicated in Figure 1.

Assessment of TOS, MDA, CAT and SOD in Serum

The serum level of the TOS, MDA, CAT and SOD are reported in Figure 1. Our results showed that CAT activity was significantly higher in case groups (Non-ICU and ICU) compared to control group: 16.34 ± 2.97, 15.54 ± 3.33 and 8.05 ± 1.72 (mU/L), respectively; *p < 0.001, (Figure 1A). CAT activity showed no significant difference between Non-ICU and ICU group. The serum level of SOD was significantly increased in case group 25.22 ± 5.75 and 34.89 ± 7 (U/mL) in comparison to control group 19.11 ± 4.72 (U/mL); *p < 0.001. Also, there was a significant difference in serum SOD level between Non-ICU and ICU group; *p < 0.01(Figure 1B). We observed that oxidative stress increases in Covid-19 patients. MDA serum levels as an indicator of lipid peroxidation, were nearly 3 times higher in case groups in comparison to control group: 1.54 ± 0.24 and 1.78 ± 0.18 (nmol/l) in Non-ICU and ICU, respectively VS 0.55 ± 0.24 (nmol/mL) in control group, *p < 0.001. Interestingly, MDA serum levels were significantly higher in ICU group compared to Non-ICU, *p < 0.05 (Figure 1C). TOS is another marker to

### Table 1. Demographic and clinical characteristics of case (ICU and non-ICU patients) and control groups

| Characteristics | Control (N = 24) | No ICU (N = 14) | ICU (N = 10) | p* |
|----------------|----------------|----------------|-------------|---|
| Age (year)     | 51.5 ± 13.58   | 57.36 ± 10.87  | 55.5 ± 15.65 | 0.402 |
| Gender         |                |                |             |    |
| Male           | 7              | 5              | 5           | 0.057 |
| Female         | 17             | 9              | 5           |      |
| Education Level|                |                |             |    |
| Illiterate     | 5              | 2              | 2           | 0.799 |
| Primary        | 12             | 6              | 3           |      |
| Diploma        | 5              | 3              | 4           |      |
| Academic       | 2              | 3              | 1           |      |
| Smoke          |                |                |             |    |
| No             | 18             | 13             | 7           | 0.369 |
| Yes            | 6              | 1              | 3           |      |
| Fever          |                |                |             |    |
| No             | -              | 5              | 0           | 0.053 |
| Yes            | -              | 9              | 10          | 100.0 |
| Cough          |                |                |             |    |
| No             | -              | 6              | 3           | 0.678 |
| Yes            | -              | 8              | 7           |      |
| Respiratory distress | | | | |
| No             | -              | 5              | 0           | 0.053 |
| Yes            | -              | 9              | 10          | 100.0 |
| SPO2           |                |                |             |    |
| ≥88            | -              | 4              | 2           | 0.506 |
| <88            | -              | 10             | 8           |      |

*p-value was assessed by exact Fisher rest.

### Table 2. The baseline characteristics of the case (ICU and non-ICU patients) and control groups

| Characteristics | Control (n = 24) | No ICU (n = 14) | ICU (n = 10) | *p |
|----------------|----------------|----------------|-------------|---|
| ESR (mm/h)     | 9.88 (7.04)    | 35.86 (14.10)  | 43.20 (17.72) | <0.001 |
| CRP (mg/l)     | 6.21 (2.21)    | 43.93 (22.80)  | 49.80 (19.14) | <0.001 |
| Ferritin (µg/l)| 90.33 (35.50)  | 517.57 (270.78) | 544.70 (188.81) | <0.001 |
| Albumin (g/dl) | 5.07 (0.48)    | 3.40 (0.38)    | 2.86 (0.97)  | <0.001 |
| White blood cell count, × 10^9 per L | 6483.33 (1632.46) | 6907.14 (4695.74) | 8110.00 (4894.99) | 0.480 |
| Neutrophil count × 10^9 per L | 64.37 (10.62) | 76.14 (13.57) | 80.80 (8.51) | <0.001 |
| Lymphocyte count × 10^9 per L | 33.46 (10.80) | 21.28 (12.31) | 16.70 (7.53) | <0.001 |
| Haemoglobin, g/l | 12.74 (2.04) | 12.61 (0.89) | 12.73 (2.35) | 0.977 |
| Platelet count × 10^9 per L | 225.92 (55.59) | 196.71 (50.49) | 217.10 (113.64) | 0.468 |

*p value was determined by ANOVA.
assess total oxidation state of the body. The mean serum concentration of TOS in Control group was 4.67 ± 1.39 (μmol H2O2 Eq/L) which was significantly less than that of in case groups 13.08 ± 1.6 in Non-ICU and 15.13 ± 2 (μmol H2O2 Eq/L), p < 0.001. There was no significant difference in TOS serum level between Non-ICU and ICU groups (Figure 1D).

Discussion

Several studies proposed the association between oxidative stress and Covid-19 pathogenesis (13,14). This study supplies proof that Covid-19 patients showed high serum level of oxidative stress and inflammatory markers and low serum level of antioxidants, in comparison with control group, especially in patients admitted in ICU. Our results presented high serum level of MDA in case groups. MDA is an important indicator of oxidative stress. There is a significant correlation between oxidative stress markers and respiratory viral infection particularly RNA viruses (15). In vitro and In vivo studies indicated that some viruses could change redox balance of cell. The beginning of oxidative stress by virus infection (such as respiratory syncytial virus) is necessary for activation of innate immunity by cytokines production (16). Besides, oxidative stress induced by several viruses involved in facilitation of virus replication inside the cell (17).

It is described the function of macrophage respiratory burst in reaction to Covid-19 infection, which can lead to ROS production (18). Over production of ROS/RNS have role in lung tissue injury and dysfunction of epithelial barrier induced by acute respiratory viral infections (19). NADPH oxidase 2(NOX2) has important role in ROS production, arterial dysfunction, and thrombosis (induced by platelet activation) (20). It is indicated NOX2 overactivation in COVID-19 patients (21).

Viruses can suppress antioxidant systems including super oxide dismutase, glutathione S-transferase, catalase, glutathione peroxidase in human alveolar type 2-like epithelial cells and small airway epithelial cells (22). Although we observed high level of antioxidant enzymes, that may be a remedial mechanism to counteract oxidative stress. CAT and SOD have major role in neutralization of free radicals including ROS and RNS (23). In line with our results, Zhu Z, et al, indicated high level of inflammatory markers such as ESR and CRP (24).

Reduced oxygen transport to the tissues, disseminated intravascular coagulopathy and sepsis are shown in
COVID-19 patients (25). Hypoxia can produce reactive species such as superoxide and H2O2 which can upregulate the expression of inflammatory cytokines (13). In turn, inflammatory cytokines can increase oxidative stress markers via activation of macrophages, neutrophils, and endothelium cells (26). These interactions between oxidative stress and inflammatory cytokines can lead to several organ failures in COVID-19 patients who proceed to worsening the condition. “Cytokine storm” in covid-19 patients is well established (27,28). Cytokine storm has prominent role in development of acute respiratory distress syndrome and various organ dysfunctions (13). Inflammasome, as a cytosolic molecular complex, senses inflammatory signals to accelerate cytokines maturation. The NLR protein NLRP3/NALP3 makes well known inflammasome (29). In fact, Inflammasome is a significant component in cytokine storm development (27). ROS as an inflammasome activator (via direct activation of NLRP3/NALP3 inflammasome) has function in inflammation induced by COVID-19 virus and following blood dissemination. As well, it is suggested that adaptive immune response to oxidative stress may be associated with systemic injury (13). Also, ROS can indirectly increase inflammasome via NF-κB activation (30,31). The results of this study indicated high serum level of antioxidants enzymes; this may be a remedial mechanism to counteract oxidative stress. CAT and SOD have major role in neutralization of free radicals including ROS and RNS (23). According to Table 3, we observed hematological disorder in COVID-19 patients: increased level of ferritin, ESR and low count of lymphocytes. Lymphopenia is reported in nearly 85% of sever Covid-19 patients (6). In covid-19, uncontrolled inflammatory responses cause low count of lymphocytes. Reduced oxygen transport to the tissues, coagulopathy and sepsis are shown in COVID-19 patients (25). Hypoxia can produce reactive species such as superoxide and H2O2 which can upregulate the expression of inflammatory cytokines (25,26). In turn, inflammatory cytokines can increase oxidative stress markers via activation of macrophages, neutrophils, and endothelium cells (8). This interaction between oxidative stress and inflammatory cytokines can lead to several organ failures in COVID-19 patients who proceed to worsening condition. Ferritin with immune-inhibitory and pro-inflammatory effects, involves in cytokine storm (32). In line with our results, Mehta P, et al, reported the high serum level of ferritin in COVID-19 patients (27). It is hypothesized that high serum ferritin level may be an influencing factor on COVID19 severity, and reduction strategies in ferritin level may decrease exacerbation of this disease (33). In this study, we observed the low level of serum albumin in COVID-19 patients. Albumin has different bioactive roles. This negative acute phase reactant is involved in extracellular antioxidant system via binding metals, scavenging free radicals, and supplying thiol group (34). In consistent with our results, Zhou F, et al. (35) and Zhang Y, et al. (36) reported low level of serum albumin in sever Covid-19 patients.

Table 3. Pairwise comparison of the characteristics of the case (ICU and non-ICU) and control groups using Tuckey post hoc

| Characteristics | Mean Difference | SD | 95% Confidence Interval | p     |
|-----------------|-----------------|----|------------------------|-------|
| ESR (mm/h)      |                 |    |                        |       |
| (control)-(Non-ICU) | -25.98 | 4.06 | -35.82 - 16.15 | <0.001 |
| (control)-(ICU) | -33.32 | 4.54 | -44.33 - 22.32 | <0.001 |
| (Non-ICU)-(ICU) | -7.34 | 5.00 | -19.45 4.77 | 0.315 |
| CRP (mg/l)      |                 |    |                        |       |
| (control)-(Non-ICU) | -37.72 | 5.05 | -49.97 - 25.47 | <0.001 |
| (control)-(ICU) | -43.59 | 5.66 | -57.30 - 29.88 | <0.001 |
| (Non-ICU)-(ICU) | -8.87 | 0.22 | -20.95 9.21 | 0.616 |
| Ferritin (µg/l) |                 |    |                        |       |
| (control)-(Non-ICU) | -427.24 | 57.23 | -565.93 - 288.55 | 0.000 |
| (control)-(ICU) | -454.37 | 64.05 | -609.59 - 299.14 | 0.000 |
| (Non-ICU)-(ICU) | -27.13 | 70.45 | -197.88 143.63 | 0.922 |
| Albumin (g/dl)  |                 |    |                        |       |
| (control)-(Non-ICU) | 1.67  | 0.20 | 1.19 2.16 | 0.000 |
| (control)-(ICU) | 2.21  | 0.22 | 1.67 2.75 | 0.000 |
| (Non-ICU)-(ICU) | 0.54  | 0.24 | -0.06 1.13 | 0.085 |
| Neutrophil count × 10⁹ per L |     |    |                        |       |
| (control)-(Non-ICU) | -11.77 | 3.77 | -20.89 - 2.64 | 0.009 |
| (control)-(ICU) | -16.42 | 4.21 | -26.64 - 6.21 | 0.001 |
| (Non-ICU)-(ICU) | -4.66 | 4.64 | -15.89 6.58 | 0.578 |
| Lymphocyte count × 10⁹ per L |     |    |                        |       |
| (control)-(Non-ICU) | 12.17 | 3.60 | 3.44 20.90 | 0.004 |
| (control)-(ICU) | 16.76 | 4.03 | 6.99 26.53 | 0.000 |
| (Non-ICU)-(ICU) | 4.59  | 4.43 | -6.16 15.33 | 0.559 |
19 patients. Huang J, et al. (37), in a retrospective study, indicated that serum level of albumin has inverse association with death risk in Covid-19 patients, independent of other markers including lymphocyte count or comorbidities. Low level of serum albumin in COVID-19 patients may be explained by systemic inflammation. The therapeutic role of albumin in inflammatory disorder (such as sepsis) is mediated via controlling effect on inflammation and oxidative stress (38,39). However, the advantage role of albumin in COVID-19 patients should be documented in more studies.

Conclusion

There is a clear association between oxidative stress and severity of several viral diseases. Our results indicated that Covid-19 patients suffer from oxidative stress which may aggravate patients, condition. One of the advantages of this study is that we exclude any underlying condition which may have effect on oxidative stress. Over production of free radicals and defect in antioxidant system have major role in SARS-COV pathogenesis. On the other hand, the cross talk between oxidative stress and cytokine storm may have significant effect on the severity of Covid-19 patient’s symptoms. It seems that strategies for reducing or preventing of oxidative stress may help in Covid-19 managements. It is noted that our study has several limitations: the first and the most important one is small sample size. Secondly, we could not collect data on dietary patterns of subjects. Thirdly, all the participants were middle-aged. More clinical studies are necessary and should address young, aged patients.

Consent to Participate

All participants provided written informed consent.

Conflict Interest

All authors declared no conflict of interest regarding this paper.

Acknowledgments

The authors appreciate the Deputy of Research and Technology, Hamadan University of Medical Sciences for their financial support in this research.

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