The effect of low level-laser irradiation on antioxidant enzymes and mineral levels in serum of patients with type 2 diabetes mellitus

Danik Martirosyan1, Mohammad Reza Ashoori2, Hossein Mirmiranpour3*

1Functional Food Center, Functional Food Institute, Dallas, TX, USA; 2Department of Laboratory Sciences, School of Health and Allied Medical Sciences, Zanjan University of Medical Sciences, Zanjan, Iran; 3Endocrinology and Metabolism Research Center (EMRC), Vali-Asr Hospital, School of Medicine, Tehran University of Medical Sciences, Tehran, Iran

*Corresponding Author: Hossein Mirmiranpour, MD, PhD, Endocrinology and Metabolism Research Center (EMRC), Vali-Asr Hospital, School of Medicine, Tehran University of Medical Science, 1th floor, Keshavarz Boulevard, Tehran, Iran.

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ABSTRACT

Background: The control and management of type 2 diabetes mellitus is the most important way to prevent health consequences of the disease. Oxidative stress derived from diabetes mellitus is an important cause of these symptoms in uncontrolled diabetic patients. The consumption of functional foods containing antioxidants and trace minerals can help prevent these consequences and control diabetes mellitus.

Objective: In this study, we examined whether low-level laser therapy could have an effect on levels of antioxidant enzymes and minerals in the serum of patients with type 2 diabetes mellitus.

Methods: Thirty individuals with type 2 diabetes and thirty healthy individuals, as controls, were selected as participants for this study. The levels of antioxidant enzymes glutathione peroxidase, catalase, superoxide dismutase; biochemical parameters, such as glucose and hydrogen peroxide; minerals iron, zinc, magnesium, copper; and selenium binding protein 1 (as an indicator of selenium) were studied before and after low level-laser therapy.

Results: The levels of antioxidant enzymes and some minerals significantly increased in control and diabetic patients after low level-laser therapy. The levels of glucose, hydrogen peroxide, and selenium binding protein 1, however, were unchanged after low-level laser therapy.

Conclusions: Low level-laser therapy may be an important tool for reducing oxidative stress caused by type 2 diabetes mellitus due to its ability to increase levels of antioxidant enzymes and mineral content.
INTRODUCTION
Diabetes mellitus (DM) is one of the most prevalent chronic metabolic diseases worldwide and is characterized by elevated levels of glucose in blood (hyperglycemia). According to the World Health Organization (WHO), 422 million people lived with diabetes in 2014 [1, 2]. Moreover, diabetes caused 1.5 million deaths in 2012 [3]. There are two main types of the disease, type 1 diabetes mellitus (T1DM) and type 2 diabetes mellitus (T2DM). T2DM results from insulin resistance or an inability to produce enough insulin. This type of diabetes is associated with poor nutrition, an unhealthy lifestyle, and low physical activity. Hyperglycemia induced by T2DM causes a rise in reactive oxygen species (ROS) production which can lead to oxidative stress and impairs the antioxidant defense system [4,5]. Oxidative stress and inflammation caused by diabetes may produce symptoms and certain health consequences of DM [6] such as skin wounds, peripheral neuropathy, nephropathy, and retinopathy [7]. The role of oxidative stress, inflammatory responses, and cell signaling have been considered in causing these consequences [8].

Functional foods have been found to be essential for the management and prevention of certain diseases such as diabetes and its sequelae [9]. Functional foods can also be sources of many micronutrients such as vitamins and minerals, which have antioxidant and anti-inflammatory properties [10]. Minerals found in functional foods are especially important for the antioxidant defense system. Antioxidants enzymes such as glutathione peroxidase (GPx), superoxide dismutase (SOD), and catalase (CAT) can be extracted from some functional foods and bioactive compounds [11-13]. These enzymes found in antioxidants are the first line of defense in the human body against oxidative stress and depend on trace minerals such as Zinc (Zn), Selenium (Se), Iron (Fe), Copper (Cu), and Magnesium (Mg) as cofactors [14]. These enzymes play a vital role in protecting the body’s biological systems against attack by free radicals [14]. It is interesting to note that there is mounting attention in investigating these antioxidants’ preventative and therapeutic abilities against disease.

Photo biomodulation, or low level-laser therapy (LLLT), has been used for more than fifty years [15]. This kind of laser, instead of producing any heat, acts through photo-chemical and photo-physical reactions [16]. Currently, LLLT appears to be an increasingly popular therapeutic option among physicians for many problems such as skin wounds and neuropathic pains [17-21]. For example, low-energy laser radiation has exerted direct and considerable influence on tissue repair processes without causing heating effects [22]. In clinical settings, LLLT has been used for accelerating healing, regeneration, and reducing pain and inflammation [23,24]. LLLT has also shown to be effective in healing skin wounds by stimulating fibroblasts and cell proliferation as well as collagen production [25]. Researchers have suggested that low level-laser therapy (LLLT) may influence oxidative stress parameters as well as alter the activity of antioxidant enzymes and output of ROS [26]. Thus, LLLT may have importance in the control of DM in the patients with T2DM.

This study reviews the effect of LLLT on the activity of several antioxidant enzymes and their trace element cofactors. The evaluation of LLLT’s effect on these minerals as a group of bioactive compounds was another goal of this investigation. A correlation between LLLT use and changes in these minerals’ quantity and alterations to above-mentioned enzymes’ function may facilitate new approaches to antioxidant activation in humans. Therefore, the aim of this study is to assess the effect of LLLT on the activity of antioxidant enzymes, serum quantity of antioxidant minerals, and related correlations between these enzymes and minerals in T2DM.

METHODS
Diagnostic kits for the antioxidant enzymes CAT and GPx were purchased from Biocore (Biocore
Diagnostik Ulm GmbH, Ulm, Deutschland). The SOD assay kit was purchased from Biovision (BioVision Incorporated, USA). The hydrogen peroxide (H₂O₂) assay kit was purchased from ZellBio (ZellBio GmbH, Ulm, Germany). Human glucose (with an intra- and inter-assay CV < 8% and < 10%, respectively) enzyme-linked immunosorbent assay (ELISA) kits were purchased from MyBioSource Inc. (San Diego, CA 92195-3308, USA). Zn, Cu, Fe, Mg and Se binding protein 1 assay kits were purchased from MyBioSource Inc. (San Diego, CA 92195-3308, USA).

Parameters were evaluated according to kit instructions. Microplate reader (Mindray, model MR-96A, Germany) and microplate spectrophotometer (model Fluostar, bmglabtech, Germany) were used for this assay. Green diode pumped solid state (DPSS) Laser Pointer (model RLP-532, 1040 Vienna, Austria) was used for LLLT.

Participants: In this research, thirty patients with T2DM and thirty healthy individuals, as a control group, were randomly selected among the persons referred to the Endocrinology and Metabolism Research Center (EMRC), Vali-Asr Hospital, Tehran University of Medical Sciences, Tehran, Iran. Written consent was obtained from healthy subjects and persons with T2DM before sampling and starting the study. People with other chronic illnesses, people undergoing surgery, and those with T1DM and infectious illness were excluded. Diagnostic criteria for T2DM was accomplished according to WHO standards for DM.

General Characteristics and Sampling: In this study, the subjects were divided into two groups: those with T2DM and healthy controls. General characteristics of participants such as sex, age, height, weight and body mass index (BMI) were recorded. Blood samples were obtained from all participants after 12 hours of overnight fasting. Blood samples were centrifuged, and their serum was separated. The obtained serum was used to evaluate the parameters described above. After 24 hours, samples were irradiated by low level green LASER diode and then examined for a second time.

Laser Irradiation: A diode laser pointer was used for irradiation. The laser was a low level green laser diode with a wavelength of 532 nm at 100 mw in a continuous wave mode with divergence < 1.5 mRad, beam mode (TEM00), beam diameter at aperture ~1.5, crystal type Nd:VYO4:KTP, and power source 1 × 3V CR2 alkaline batteries. The power density was 509.55 mW/cm² at a distance of 6.5 cm from the laser device from serum inside the tube, and the diameter of the laser spot was set to 0.5 cm. Irradiation was applied for 8 seconds.

Statistical Analysis: Statistical analysis was accomplished utilizing SPSS (version 23, IBM, USA) software for Windows. All results were expressed as mean ± standard deviation. Paired sample t-test and independent sample t-test were used to compare the mean of the obtained data. In all analyses, a p-value of <0.05 qualified the presence of a statistically substantial difference.

RESULTS

General Characteristics: The age of participants in this study was between 55 and 75 years old. Among healthy controls, 60% were female and 40% were male. 37% of people with T2DM were female and 63% of people with T2DM were male. General characteristics between the control group and diabetic group are shown in Table 1. In the diabetic group, height, weight, and body mass index (BMI) were not significant in comparison with the healthy control group.

Table 1. Participant’s General Characteristics

<table>
<thead>
<tr>
<th>Characteristic</th>
<th>Control Group n = 30</th>
<th>Diabetic Group n = 30</th>
<th>P value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Height (Cm)</td>
<td>165.3±5.6</td>
<td>165.8±6.6</td>
<td>0.72</td>
</tr>
<tr>
<td>Weight (Kg)</td>
<td>78.2±6.2</td>
<td>79.9±8.0</td>
<td>0.35</td>
</tr>
<tr>
<td>BMI (Kg/m²)</td>
<td>28.6±2.1</td>
<td>29.0±1.5</td>
<td>0.47</td>
</tr>
</tbody>
</table>

Data are given as mean ± SD. P value < 0.05 is significant.

BMI, body mass index

Biochemical Parameters: The levels of antioxidant enzymes CAT, SOD, and GPx were quantified in the collected samples. The levels of aforementioned minerals associated with these enzymes as well as glucose and H₂O₂ were also measured. The
concentrations of these parameters were measured before and after LLLT (Table 2) and compared in the control and diabetic groups. The results of this study are shown in Table 3.

As seen in Table 2, there was a significant difference in the concentrations of catalase, GPx, and SOD in the control and diabetic groups before and after LLLT (P value < 0.001). Comparison of glucose, H₂O₂, and Sebp1 concentrations did not show any significant difference in the control and diabetic groups before and after LLLT (P value > 0.05). There was also no significant difference in the comparison of concentrations of minerals (Fe, Zn, Mg and Cu) in the control group before and after LLLT (P value > 0.05). However, there was a significant difference in the concentration of these minerals in the diabetic group before and after LLLT (P value was 0.01, 0.005, 0.001, and 0.001, respectively for Fe, Cu, Mg and Zn). The comparison of mean concentration between all biochemical parameters in the control and diabetic groups after LLLT showed a significant difference (P value < 0.001).

The relationship between the intake of these bioactive compounds, diabetes, and LLLT as well as changes in the concentration of antioxidant enzymes and minerals is summarized in Figure 1.

![Diagram](https://via.placeholder.com/150)

**Figure 1.** The relationship between diabetes, antioxidants, and LLLT. (A) The relationship between functional foods intake, antioxidants, and reduction of oxidative stress formation by inhibition of reactive oxygen species. (B) The effect of LLLT on antioxidant enzymes and some minerals in the serum of diabetic patients before and after LLLT.
DISCUSSION

In this study, we aimed to investigate the effects of LLLT on the quantity of several biochemical parameters in patients with T2DM. The therapeutic effect of LLLT on diabetic patients has been evaluated in many studies. In one study by Denadai et al., it was reported that LLLT (660 nm) was effective in reducing levels of oxidative stress in diabetic rats [27]. In one of our previous studies, the effect of LLLT on the activity of CAT was investigated. In that study, we found that green (530nm) and blue (450nm) laser light increased CAT activity in blood samples more than red laser light [28].

In terms of increased CAT activity, this study agrees with our previous study. Lim et al. also concluded that LLLT (670 nm) increased CAT activity and expression. In their experiments, LLLT had no effect on GPx and SOD activity. [29] In this study, no reduction in H2O2 by LLLT was observed in the control and diabetic groups. However, GPx and SOD concentrations increased significantly in both the control and diabetic groups. By reducing oxidative stress through increasing the activity of antioxidant enzymes, such as CAT and SOD, health consequences of diabetes may be reduced. LLLT may be an effective method to increase the activity of antioxidant enzymes, according to previous studies and the results of this study.

In our study, LLLT had no effect on lowering blood glucose, which also agrees with the study by Lim et al. However, Longo et al. reported that LLLT is effective in lowering blood glucose for patients with T1DM [30].

According to the new definition of functional foods by the Functional Foods Center (FFC) [31], any functional food product contains one or more biologically active compounds such as minerals, vitamins, and antioxidant enzymes. The therapeutic effects of some dietary supplements containing such biologically active compounds, which have anti-

<table>
<thead>
<tr>
<th>Parameters</th>
<th>Control Group n = 30</th>
<th>Diabetic Group n = 30</th>
</tr>
</thead>
<tbody>
<tr>
<td>CAT (U/ml)</td>
<td>Before LLLT: 2.7±0.07 After LLLT: 2.9±0.03 P value: &lt; 0.001</td>
<td>Before LLLT: 1.5±0.2 After LLLT: 2.6±0.2 P value: &lt; 0.001</td>
</tr>
<tr>
<td>GPx (U/ml)</td>
<td>121.8±2.3 Before LLLT: 122.7±2.4 P value: &lt; 0.001</td>
<td>81.5±6.2 Before LLLT: 83.8±6.3 P value: &lt; 0.001</td>
</tr>
<tr>
<td>SOD (U/ml)</td>
<td>7.6±0.3 Before LLLT: 7.8±0.3 P value: &lt; 0.001</td>
<td>4.0±0.3 Before LLLT: 4.8±0.4 P value: &lt; 0.001</td>
</tr>
<tr>
<td>Glucose (μg/ml)</td>
<td>302.6±19.5 Before LLLT: 301.8±19.0 P value: 0.86</td>
<td>424±23.1 Before LLLT: 420±20.9 P value: 0.57</td>
</tr>
<tr>
<td>H2O2 (μM/ml)</td>
<td>223.6±19.6 Before LLLT: 221.8±19.6 P value: 0.71</td>
<td>332.3±27.5 Before LLLT: 323±23.0 P value: 0.23</td>
</tr>
<tr>
<td>Sebp1 (pg/ml)</td>
<td>4971.1±289.6 Before LLLT: 4976.1±289.6 P value: 0.95</td>
<td>3171.1±289.6 Before LLLT: 3221.2±289.8 P value: 0.54</td>
</tr>
<tr>
<td>Fe (μmol/l)</td>
<td>292.3±11.3 Before LLLT: 291.3±11.2 P value: 0.74</td>
<td>397.6±14.3 Before LLLT: 387.3±14.4 P value: 0.01</td>
</tr>
<tr>
<td>Cu (μmol/l)</td>
<td>295.3±12.3 Before LLLT: 294.2±12.2 P value: 0.66</td>
<td>399.4±13.9 Before LLLT: 389.3±13.9 P value: 0.005</td>
</tr>
<tr>
<td>Mg (mmol/l)</td>
<td>7.0±0.8 Before LLLT: 7.1±0.7 P value: 0.47</td>
<td>3.9±0.8 Before LLLT: 4.7±0.9 P value: 0.001</td>
</tr>
<tr>
<td>Zn (μmol/l)</td>
<td>155.8±6.6 Before LLLT: 156.9±6.5 P value: 0.51</td>
<td>99.1±5.9 Before LLLT: 105.2±5.9 P value: 0.001</td>
</tr>
</tbody>
</table>

Results are showed as mean ± SD. P value < 0.05 is significant. LLLT, low level-laser therapy; CAT, catalase; GPx, glutathione peroxidase; SOD, superoxide dismutase; Sebp1, selenium binding protein 1’ Fe. Iron; Cu, Copper; Mg, Magnesium; Zn, Zinc.
Table 3. Comparison between the concentrations of biochemical parameters after LLLT in two groups.

<table>
<thead>
<tr>
<th>Parameters</th>
<th>Control Group n = 30</th>
<th>Diabetic Group n = 30</th>
<th>P value</th>
</tr>
</thead>
<tbody>
<tr>
<td>CAT (U/ml)</td>
<td>2.9±0.03</td>
<td>2.6±0.2</td>
<td>&lt; 0.001</td>
</tr>
<tr>
<td>GPx (U/ml)</td>
<td>122.7±2.4</td>
<td>83.8±6.3</td>
<td>&lt; 0.001</td>
</tr>
<tr>
<td>SOD (U/ml)</td>
<td>7.8±0.3</td>
<td>4.8±0.4</td>
<td>&lt; 0.001</td>
</tr>
<tr>
<td>Glucose (μg/ml)</td>
<td>301.8±19.0</td>
<td>420±20.9</td>
<td>&lt; 0.001</td>
</tr>
<tr>
<td>H2O2 (μM/ml)</td>
<td>221.8±19.6</td>
<td>323±23.0</td>
<td>&lt; 0.001</td>
</tr>
<tr>
<td>Sebp1 (pg/ml)</td>
<td>4976.1±289.6</td>
<td>3221.2±289.8</td>
<td>&lt; 0.001</td>
</tr>
<tr>
<td>Fe (μmol/l)</td>
<td>291.3±11.2</td>
<td>387.3±14.4</td>
<td>&lt; 0.001</td>
</tr>
<tr>
<td>Cu (μmol/l)</td>
<td>294.2±12.2</td>
<td>389.3±13.9</td>
<td>&lt; 0.001</td>
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<td>Mg (mmol/l)</td>
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Results are showed as mean ± SD. P value < 0.05 is significant. LLLT, low level-laser therapy; CAT, catalase; GPx, glutathione peroxidase; SOD, superoxide dismutase; Sebp1, selenium binding protein 1; Fe, Iron; Cu, Copper; Mg, Magnesium; Zn, Zinc.

Inflammatory and antioxidant properties have been discussed [32]. Thus, functional foods containing such compounds may be effective in managing diabetes and other chronic diseases [34]. Zn, which could be considered as a bioactive compound in functional food, is one of the essential trace elements in humans and Zn deficiency has been observed in diseases such as DM [35]. Zn acts as an insulin mimetic in the insulin signal transduction pathway and therefore has insulin-like effects in glucose homeostasis [36]. In one study, Abdel-Magied et al. investigated the effect of LLLT (870 nm) on certain minerals and enzymes in the liver and kidney of rats. They reported that LLLT increased the activity of antioxidant enzymes such as SOD and CAT as well as the abundance of minerals such as Fe, Cu, Zn, Mg, and Se [37]. Selenium can act as an anti-inflammatory and antioxidant factor, and acts as a cofactor in selenoproteins such as SOD [38]. We studied Sebp1 as an indicator of Se concentration. In our study LLLT significantly increased levels of Zn and Mg in diabetic patients, but decreased levels of Cu and Fe. The concentration of Sebp1 was not significantly different in diabetic and control subjects before and after LLLT.

However, the difference in levels of certain minerals and enzymes was significant between control and diabetic subjects after LLLT. In a study by Atalay et al. [38] where several trace elements were studied in the serum of women with T2DM, it was suggested that levels of minerals such as Mg, Fe, Cu, and Zn were low in women with T2DM. These differences in mineral levels between diabetic and non-diabetic individuals may express the relationship between glucose metabolism and these minerals. Low Mg levels, for example, are associated with poor glycemic control [39]. According to the above results and the results of our study, the activity of antioxidant enzymes as well as trace elements should be considered in diabetic patients. LLLT and the consumption of functional foods containing antioxidant enzymes and trace minerals may play an important role in increasing the levels of these parameters.

**CONCLUSION**

Our study showed that LLLT use in patients with T2DM may be used to increase levels of antioxidant enzymes and minerals and alleviate diabetes-
associated oxidative stress. Intake of bioactive compounds such as Mg, Zn, Se, Fe, and Cu through functional foods may also play an important role in controlling DM and preventing its health consequences.

LIST OF ABBREVIATIONS
LLLT, low level-laser therapy; DM, diabetes mellitus; T1DM, type 1 diabetes mellitus; T2DM, type 2 diabetes mellitus; WHO, world health organization; GPx, glutathione peroxidase; SOD, superoxide dismutase; CAT, catalase; ROS, reactive oxygen species; BMI, body mass index; Sebp1, selenium binding protein 1; Fe, Iron; Zn, Zinc; Mg, Magnesium; Cu, Copper.

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COMPETING INTERESTS
The authors declare that there are no conflicts of interest.

AUTHOR’S CONTRIBUTIONS
Danik Martirosyan participated in the study design. He also edited the article. Mohammad Reza Ashoori participated in the writing and analysis of the results. Hossein Mirmiranpour contributed to the original idea of the paper, doing the experimental work and, data collection. All authors read and approved the final version before its submission.

HUMAN AND ANIMAL STUDIES
This article contains human studies. All of which were consented and humane. No animal studies were conducted.

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