UV-vis spectroscopic study of lipid peroxidation and antioxidant contents of kidney of Albino mice co-administrated with sildenafil citrate and alcohol

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Abstract. Male Wistar Albino mice were co-administrated with appropriate dose of sildenafil citrate and alcohol. The kidney samples of those animals were analyzed for lipid peroxidation and antioxidant contents. Lipid peroxidation was measured based on TBARS. Antioxidant such as SOD, CAT, GPx and GSH were also determined. From the present investigation, an enhanced level of TBARS accompanied by a decreased concentration of enzymic as well as non-enzymic antioxidants in kidney was detected for 30 days animals treated concomitantly with sildenafil citrate and alcohol. Increased level of lipid peroxidation lead to accumulation of free radicals and increased oxidative threat in kidney tissues. Also, the decreased concentration of antioxidant enzymes in renal tissues indicates the failure of antioxidant defence system.

Keywords: Albino mice, kidney, sildenafil citrate, alcohol, lipid peroxidation, antioxidants

1. Introduction

The issue of erectile dysfunction provides a paradoxial situation to both the patients and physicians. Sildenafil citrate is reportedly an effective and safe medication [2] indicated for the treatment of ED. It is a competitive inhibitor of cGMP-Specific phosphodiesterase type 5. The medication amplifies the effects of sexual stimulation by regarding the degradation of this enzyme. Sildenafil has been found effective in several subpopulations of men with ED, including sufferers from diabetes, hypertension, spinal cord injuries [7], multiple sclerosis [11], depression [10,36], PTSD [25], men after resection of the prostate or radical prostatectomy [22], after renal transplant [30], men on dialysis [5] and men aged 65 years and older [3].

Studies conducted by Jeffcoate et al. [15] and McCambridge et al. [19] show that men with ED are frequently chronic alcohol addicts. The findings of previous studies show that modest ethanol doses (e.g., at blood concentrations of \( \leq 100 \) mg/dl) can both increase sexual drive and decrease erectile capacity in men [4]. As a result, alcohol dependent men commonly suffer from erectile dysfunction being an effective vasoactive agent currently available for the treatment of ED, most of them uses sildenafil citrate.
The combined use of sildenafil and alcohol may affect the biochemical balance of the body. In human body, kidneys are the organs primarily responsible for regulating the amounts and concentration of lipid peroxidative and antioxidant substances in extra cellular fluid. Hence, here, we choose the experimental organ as kidney.

Previous studies reveal that alcohol consumption leads to kidney swelling and reduced kidney function [37] and also, kidney cells get enlarged with increased amounts of protein, fat and water. Alcohol consumption reduces the amount of potassium excreted by the kidneys. Excessive alcohol consumption can have profound negative effects on the kidneys and their function in maintaining the body’s fluid, electrolyte and acid–base balance, leaving alcoholic people vulnerable to a host of kidney related health problems [21]. These points must be kept in our mind during the analysis of this work.

The goal of the present study was whether the combined consumption of sildenafil citrate and alcohol affects the lipid peroxidation and antioxidant levels in the kidney tissues. We sought to examine (i) the concentration of lipid peroxidative substance such as TBARS, and (ii) the concentration of antioxidants such as SOD, CAT, GPx and GSH before and after the co-administration of sildenafil and alcohol.

2. Materials and methods

2.1. Chemicals

Drug, here, refers to the commercially available 50 mg tablet of sildenafil citrate (VIAGRA). Alcohol was purchased from sigma chemical Co. (St Louis, MO, USA). All other chemicals utilized were of analytical grade and were obtained from local firms (India).

2.2. Animals

The local institutional animal ethics committee (Register number 160/1999/CPCSEA), Annamalai University, Annamalai Nagar, India, approved the experimental design. The animals were maintained as per the principles and guidelines of the ethical committee for animal care of Annamalai University in accordance with Indian National Law on animal care and use. Healthy male Wistar Albino mice (Mus musculus), with an initial body weight of 25–30 g, were used in this study. The animals were housed in stainless steel mesh cages, housed under controlled conditions (temperature 25 ± 2°C, natural high-dark cycle). Commercial standard pellet diet (Hindustan lever Ltd, Mumbai, India) and drinking water were provided ab libitum. Six animals were usually tested as a group in each experiment. The commercial mice feed contained 5% fat, 21% protein, 55% nitrogen free extract and 4% fiber (w/w), with adequate minerals and vitamin contents.

2.3. Experimental design

The animals were randomly divided into seven groups of six animals each.

Groups S1: Control animals treated intragastrically with conductivity water (1 lig/gm body wt/day) for 30 days.
Group S2: Animals received drug orally (at 1 µg/gm body wt/day) for 15 days using intragastric tube.
Group S3: Animals received drug (at 1 µg/gm body wt/day administrated orally) for 30 days.
Group S4: Animals were treated with alcohol orally (at 0.01 µg/gm body wt/day) for 15 days using intragastric tube.
Group S5: Animals were treated with alcohol orally (at 0.01 µg/gm body wt/day) for 30 days.
Group S6: Animals received drug (at 1 µg/gm body wt/day) followed by oral administration of alcohol (at 0.01 µg/gm body wt/day) for 15 days.
Group S7: Animals received drug (at 1 µg/gm body wt/day) followed by oral administration of alcohol (at 0.01 µg/gm body wt/day) for 30 days.

After four hours of drug administration, the animals were sacrificed by cervical decapitation. The kidney was dissected out and quickly rinsed in 4% saline. Tissue samples were dried and homogenized using appropriate buffer by Tefflon Pestle. The following parameters were estimated using UV-vis spectrophotometer.

<table>
<thead>
<tr>
<th>Sl no.</th>
<th>Biochemical parameter</th>
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<tbody>
<tr>
<td>1</td>
<td>TBARS</td>
</tr>
<tr>
<td>2</td>
<td>GSH</td>
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<tr>
<td>3</td>
<td>GPx</td>
</tr>
<tr>
<td>4</td>
<td>SOD</td>
</tr>
<tr>
<td>5</td>
<td>CAT</td>
</tr>
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</table>

Lipid peroxidation in kidney was estimated by measuring thiobarbituric acid reactive substances (TBARS). TBARS was estimated using the method of Nichans and Samuelson [24] and reduced glutathione (GSH) by the method of Ellman [9]. Catalase (CAT) was assayed using the method of Sinha [31] and GPx by the method of Rotruck et al. [29]. Superoxide dismutase (SOD) activity was determined by the modified method of NADH-phenazinemetho sulphate-nitroblue tetrazolicem formazon inhibitor reaction, measured spectro-photometrically at 560 nm [16].

2.4. Statistical analysis

All data were expressed as mean ± SD of number of experiments (n = 6). The statistical significance was evaluated by one-way analysis of variance (ANOVA) using SPSS version 9.0 (SPSS, Cary, NC, USA) and the individual comparisons were obtained by Duncan’s Multiple Range Test (DMRT). P < 0.05 was considered to be statistically significant.

3. Results

Table 1 portrays the results of the level of lipid peroxidation in terms of TBARS and the concentration of antioxidants in terms of SOD, CAT, GPx and GSH, as carried out in the present study. Figure 1 illustrates the graphical representation of the level of TBARS for each sample. An enhanced level of TBARS in kidney has been noticed for all the experimental animals. However, statistically significant increase in TBARS concentration has been observed only for S3 (30 days drug treated), S5 (30 days alcohol treated), S6 (15 days ‘drug + alcohol’ treated) and S7 (30 days ‘drug + alcohol’ treated) groups of animals. The combined dosage of drug and alcohol administered continuously for 30 days, resulted in an elevated lipid peroxidation in kidney of Albino mice (Fig. 1).

Changes in SOD experiment have been detected in the case of kidney of Albino mice treated with the drug and alcohol, both individually and combinatorially. The decrement in SOD content has been observed to be statistically significant for S3 (30 days drug treated), S5 (30 days alcohol treated), S6 (15 days ‘drug + alcohol’ treated) and S7 (30 days ‘drug + alcohol’ treated) groups of animals (Fig. 2).
Table 1

Thiobarbituric acid reactive substances (TBARS) and antioxidants content in kidney of Albino mice treated combinatorially with sildenafil citrate and alcohol

<table>
<thead>
<tr>
<th>Group</th>
<th>Concentration of TBARS (nmol/g tissue)</th>
<th>Concentration of antioxidants</th>
</tr>
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<tbody>
<tr>
<td></td>
<td></td>
<td>SOD (unit/min/mg protein)</td>
</tr>
<tr>
<td>S1</td>
<td>1.56 ± 0.11a</td>
<td>11.06 ± 0.77a</td>
</tr>
<tr>
<td>S2</td>
<td>1.79 ± 0.11a</td>
<td>10.80 ± 0.80a</td>
</tr>
<tr>
<td>S3</td>
<td>2.45 ± 0.19b</td>
<td>10.05 ± 0.72b</td>
</tr>
<tr>
<td>S4</td>
<td>1.69 ± 0.13a</td>
<td>10.31 ± 0.79a</td>
</tr>
<tr>
<td>S5</td>
<td>2.03 ± 0.14b</td>
<td>10.92 ± 0.76b</td>
</tr>
<tr>
<td>S6</td>
<td>2.12 ± 0.11b</td>
<td>10.56 ± 0.70b</td>
</tr>
<tr>
<td>S7</td>
<td>2.67 ± 0.20f</td>
<td>9.33 ± 0.73c</td>
</tr>
</tbody>
</table>

Notes: Values are mean ± SD of six mice from each group. Values not sharing a common superscript letter differ significantly at P < 0.05 (DMRT), P < 0.05 (ANOVA). TBARS – thiobarbituric acid reactive substances; SOD – superoxide dismutase; CAT – catalase; GPx – glutathione peroxidase; GSH – reduced glutathione. S1 – control; S2 – drug (15 days); S3 – drug (30 days); S4 – alcohol (15 days); S5 – alcohol (30 days); S6 – drug + alcohol (15 days); S7 – drug + alcohol (30 days).

Fig. 1. Change in the concentration of thiobarbituric acid reactive substances (TBARS) in kidney of control and experimental Albino mice. S1 – control; S2 – drug (15 days); S3 – drug (30 days); S4 – alcohol (15 days); S5 – alcohol (30 days); S6 – drug + alcohol (15 days); S7 – drug + alcohol (30 days).

Prominent variations in the concentration of CAT have been noticed for the kidney samples of all the experimental Albino mice, compared to the control ones. The level of CAT has been found to decrease with the duration of the experiment and it is the least for S7 (30 days ‘drug + alcohol’ treated) group of animals (Fig. 3).
Fig. 2. Change in the concentration of superoxide dismutase (SOD) in kidney of control and experimental Albino mice. S1 – control; S2 – drug (15 days); S3 – drug (30 days); S4 – alcohol (15 days); S5 – alcohol (30 days); S6 – drug + alcohol (15 days); S7 – drug + alcohol (30 days).

Fig. 3. Change in the concentration of catalase (CAT) in kidney of control and experimental Albino mice. S1 – control; S2 – drug (15 days); S3 – drug (30 days); S4 – alcohol (15 days); S5 – alcohol (30 days); S6 – drug + alcohol (15 days); S7 – drug + alcohol (30 days).
A close scrutiny of Table 1 indicates the fact that the level of GPx had fallen significantly with the duration of the drug treatment. The concomitant administration of the drug and alcohol to Albino mice culminates in rapid decrease in the concentration of GPx, as compared to S1 (control) group of animals (Fig. 4).

The data presented in Table 1 show that the continuous treatment of drug to Albino mice for 15 days did not produce any tangible alterations in GSH environment while the 30 days drug treatment resulted in a statistically significant fall in it. Similar trend has also been noticed for alcohol treatment. However, the combined dosage of drug and alcohol to Albino mice end up in significant decrease in GSH level in kidney of these animals (Fig. 5).

4. Discussion

From the present study, it is well understood that the combined administration of sildenafil citrate and alcohol induced lipid peroxidation (LP) as well as had an inhibitory effect on the antioxidant enzymes. Membrane lipid peroxidation plays an important role in cell physiology and pathology and the products formed during this process have been noticed to produced unfavorable effects locally as well as far from their site of formation [28]. According to Mayya et al. [26], LP is a destructive process and the toxic products formed during it are more responsible for the membrane damage. The participation of these products in membrane fusion has been reported by De Duve et al. [6]. The disorders in lipid organization of biological membranes result in alterations in the activity of a number of membrane bound enzymes [13]. A close correlation between the membrane lipid peroxidation and some of the cell functions as well as many pathological processes has been observed by Eduards et al. [8].
Increases in lipid peroxidation leads to cell damage. According to Karine et al. [1], once lipid peroxidation is initiated, it can easily be propagated in the cellular environment where unsaturated lipids and oxygen are present. The oxidation of lipids is propagated by radical mediated chain reactions. Malon dialdehyde (MDA) and a wide range of other oxidation products are formed due to these chemical processes. Formation of MDA and other aldehydes has been regarded as significant because these compounds are toxic, mutagenic, tumorigenic and genotoxic [34]. Powerful damage to subcellular structures has occurred due to LP [26].

Co-administration of sildenafil and alcohol induces lipid peroxidation that could alter the structure and function of the cellular membrane and block cellular metabolism leading to cytotoxicity, i.e., possibility to inhibition of synthesis of cellular macromolecules.

Lipid peroxidation affects plasma membrane integrity as well as that of the nucleus and other cellular organelles lipid peroxidation may also be related to the disturbances of cell signaling processes, genotoxicity, mutagenicity and tumor promotion.

Enhanced lipid peroxidation constitutes a complex chain reaction of free radicals, while leads to a degradation of polyunsaturated fatty acid in cell membranes [12]. Free radicals cause cell injury by damaging lipids, protein and DNA if generated in excess or else antioxidant defence mechanism is impaired and may ultimately result in damage to many systems in the body. The lipid composition determines the fluidity, stability and permeability of membranes and there by influences the function of enzymes, ion channels and receptors.

In the present investigation, high levels of lipid peroxidation were detected in kidney of Albino mice treated with sildenafil and alcohol. The imbalance between production and elimination of free radicals
Sildenafil citrate and alcohol disrupt the antioxidant balance of tissues that leads to biochemical and physiological dysfunction [23]. Peroxidation of lipids, implicated in wide variety of physiological events, is a complex process that occurs during innumerable pathological condition [17]. Peroxide formation in vivo many produce serious consequences in the tissues such as oxidative destruction of thiol groups of amino acids and proteins [18]. Lipids, particularly unsaturated fatty acids, are damaged directly by free radical oxidation. Peroxidative damages induced in the cell are encountered by the elaborate defence mechanisms including enzymic and non-enzymic antioxidants [14].

Numerous studies demonstrate that changes in the antioxidant system and impairment of sodium and potassium transport are essential factors involved in the kidney tissues [32]. Evidences suggest that the excessive production of free radicals and the imbalance between oxidative species and antioxidant defence is related to the pathogenesis of neurodegenerative diseases [39].

The significant decrease in the levels of non-enzymic antioxidants, as observed in the present study, could lead to increased susceptibility of the tissue to free radical damage. The decreased concentration of antioxidant enzymes were observed in the renal tissue of the present work, which indicate the failure of antioxidant defence system. Thus, the inhibition of enzymes involved in free radical removal leads to the accumulation of H$_2$O$_2$, which promotes lipid peroxidation and modulation of DNA altered gene expression and cell death [33,38].

SOD constitutes an important link in the biological defence mechanism through disposition of endogenous cytotoxic superoxide radicals that are deleterious to polyunsaturated fatty acids (PUFA) and structural proteins of plasma membrane. So, decrease in concentration of SOD, due to the combined dosage of drug and alcohol will adversely affect the biological defence mechanism. The inhibition of CAT and GPx results in excessive accumulation of H$_2$O$_2$ in the mitochondria and peroxisomes or cytosol or both [20]. According to Rister et al. [27], increased lipid peroxidation may result in the accumulation of H$_2$O$_2$, thereby CAT activity decreases and hence, it may result in oxidative stress.

GSH plays a key role in cellular defence against toxic chemicals. It is well established that the non-enzymic antioxidants such as vitamin C and vitamin E concomitantly decreased with GSH [35]. GSH is the most abundant non-protein thiol that maintains the cellular redox status and providing first line of antioxidant protection against oxidative stress. So, low concentration of GSH due to the concomitant administration of drug and alcohol, as noticed in the present work, may reversely affect the oxidant defence.

Hence, it is concluded from the present investigation that the decreased activities of antioxidant systems and the increased level of lipid peroxidation may ultimately lead to the accumulation of free radicals and increased oxidative threat in tissue.

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