HYPOCHOLESTEROLEMIC EFFECT OF MICROWAVE ASSISTED DEFATTED FLAXSEED EXTRACT IN EXPERIMENTAL RATS

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ABSTRACT

The research was aimed to determine the hypocholesterolemic property of microwave assisted defatted flaxseed extract (MADFEs) on rat model under study. The study trial included 25 male rats which were separated into 5 groups having 5 rats in each group. Twenty rats out of 25 were hypercholesterolemic and remaining 5 were normal rats. High cholesterol diet was administered for 15 days in order to induce hypercholesterolemia, and after the induction these rats were administered different concentrations of MADFEs (150, 300 & 450mg/kg B.W.) for a time interval of 28 days. A significant reduction was observed in the levels of lipid profile parameters by the administration of extracts. The extract at a dose of 300mg/kg B.W. was shown to have provided optimal results displaying lowered cholesterol content, triglyceride and LDL-c by 16.79%, 15.41% and 12.68%, while the level of HDL elevated by 14.71% in comparison to the control F group. Likewise, the percent increase in the concentrations of catalase and SOD by 10.89% and 29.38%. The study determined that MADFEs displayed hypocholesterolemic effect and can be utilized for its therapeutic benefits. The lignin present in defatted flaxseed provide with the potential against diseases.

Keywords: defatted flaxseed, hypercholesterolemia, SDG lignin, microwave assisted extract, phenolic, enterodiol

INTRODUCTION

Hyperlipidemia is termed as elevated serum total cholesterol, triglycerides, LDL and VLDL levels, along with decrease in HDL levels is considered to be good cholesterol (Rahaman et al., 2013). It is hazardous for cardiovascular diseases, consisting of stroke and coronary heart disease. CVD is the cause of mortality around the globe that accounts around 30% of deaths per year worldwide (Naik et al., 2018). Flaxseeds as a whole are the source of lignins, soluble and insoluble fibers, omega -3 fatty acids. Main lignin found in flaxseed, mostly present in defatted flaxseed is secoisolariciresinol diglucoside (SDG). It is known to have great antioxidant, anti-inflammatory, angiogenic and anti-apoptotic properties. Among valuable benefits of defatted flaxseed extract, its oil is an optimum source of fatty acids (omega -3)/polyunsaturated fatty acids. Defatted flaxseed cake is obtained after oil extraction and used as a flaxseed meal by grinding it (Aslam et al., 2019). Natural products are getting fame in terms of less toxicity worldwide, and defatted flaxseed is a valuable by-product after oil extraction that can be used to treat diseases instead of wasting it. The flaxseed fiber reduces the levels of blood cholesterol by halting and decreasing their absorption from intestine (Zauwar et al., 2013). Recently, there is a huge interest in nutraceuticals and functional food, and many studies have been shown that etiology of many degenerative diseases along with hepatopathies and organ damage is due to reactive oxygen species (ROS) and oxygen free radicals (Rajesh et al., 2006). Ferulic acid, gallic acid, caffeic acid, vanillic acid, p-coumaric acid, sinapic acid and syringic acid are the phenolics found in the flaxseed. Lignans possess a number of biological activities i.e. antitumor activity, cardiovascular, neuro- protection, immunity, antiviral and neurotropic actions (Aslam et al., 2019). Flaxseed lignans that are not steroid phytoestrogens had structural similarity to that of leaf estrogens, which suggests that mammals exhibit estrogenic effects. Intestinal bacteria metabolize flax tissue for bioavailability in plasma (Mali et al., 2019). The SDG content varies between 9and 30 mg g-1 of defatted flaxseed meal. Phenolic acids and its derivatives in flax varied between 8-10 mg g-1 of seeds (Waszkowiak et al., 2015). Flax lignans act as Phytoestrogens and antioxidants both. Linseed contains about 800 times greater lignans than other plant foods. The SDG present in foods and flaxseeds converted into enterocolonol by gut bacteria that provide weak estrogenic or anti-estrogenic effects along with antioxidant effects. Consumption of flaxseed signifies its multiple benefits to health including reduced tumor growth, lowering serum cholesterol concentrations and lowered incidence of breast, prostate, and colon cancers (Russo and Reggiani, 2015). Flaxseed cake is most concerned source of lignin as compared to whole flaxseed (The et al., 2015). Extraction of SDG by conventional methods results in less extraction yield and it is time consuming. Therefore, nowadays, latest techniques are being used owing their efficient yield and rapid nature. Microwave assisted extraction MAE is used by many scientists for extraction of defatted flaxseed and its phenolics. MAE has recently been used for the extraction of various bioactive molecules from tea, citrus and grapes (Abdi et al., 2017). In view of all the concerned health benefits, present study is conducted to analyze the lipid lowering effects of defatted flaxseed extract by microwave assisted technology and antioxidant indices in experimental rats.
MATERIAL AND METHODS

Chemicals and reagents

All the chemicals and reagents used in the study were purchased from Sigma-Aldrich Corporation (USA). All of the chemicals used in the study were of analytical grade.

Plant material collection and authentication

Flaxseeds were purchased from local market of Lahore Pakistan. Flaxseeds were milled to reduce their particle size. Ground flaxseeds (50 g) were defatted by following the procedure described by AOAC (Horwitz, 2006) through Soxhlet apparatus (45-55°C) using n-hexane (200-250 mL) as a solvent for 5-7 h. For the removal of residual solvents from defatted flaxseeds they were placed in hot air oven (40°C). Oil and defatted flaxseeds were stored separately at 4 °C till further analysis.

MAE extraction procedure

Defatted flaxseeds of each mentioned variety was subjected to microwave assisted extraction using a Haier microwave system (Haier Corporation, Qingdao, China) operating at variable power i.e. 50 to 300 W. Defatted flaxseed (1 g) of each variety was placed in a quart tube and added up with 25 mL of methanol (70%) and 1 M NaOH. MAE for each sample was carried out at 150 W (power) for 5 minutes (time). Afterwards, extracts were treated with acetic acid for neutralization purpose and centrifuged for collection of supernatant. Before antioxidant characterization and quantification of major phenolics in MAE defatted flaxseed extracts of each variety, the supernatant was filtered (0.45 µm).

Experimental animals

For rat bioassay, 25 male albino rats were housed in animal room of IMBB. The University of Lahore. The rats were acclimatized by feeding on basal diet for a period of one week. The environmental conditions were controlled through the trial like temperature (23 ± 2 °C) and relative humidity (55 ± 5%) along with 12-h light-dark period. The experimental protocol was based on the institutional guidelines for the care and use of laboratory animals, University institute of molecular biology and biotechnology (IMBB), the University of Lahore, Lahore Pakistan. The study was approved by ethical committee of the department. Experiments on animals were performed in the animal house of IMBB, the University of Lahore.

Sample size

25 male albino rats (05 in normal group; 05 in hypercholesterolemic control group and 15 in hypercholesterolemic experimental groups were used in the experiment for rat bioassay. Rats were housed in the Institute of Molecular Biology and Biotechnology (IMBB) animal room to assess the effect of defatted flax seed extract.

Sampling technique

Rats was anesthetized by isoflurane and samples were collected by cardiac puncture at following intervals; 0 day (to establish baseline trend), 15th day (to authenticate the induction of hypercholesterolemia) and then on 21st day post administration of extract (to validate the effect of mentioned treatments).

Grouping and experimental treatment plan

Study was divided into 5 groups i.e. N0, F0, F1, F2, and F3. Group N0 was subjected to normal basal diet throughout the study, while group F0 was subjected to hypercholesterolemic diet without the administration of any extract. On the other hand, remaining groups i.e. F1, F2 and F3 was feed on hypercholesterolemic diet along with various concentrations (150mg/kg B. W. 300mg/kg B. W and 450mg/kg B. W) of extracts. Group F0, F1, F2, & F3 was subjected to high cholesterol diet for first 15 days for induction of hypercholesterolemia. All the respective extracts were administrated to each rat by oral gavage along with high cholesterol diet. In this rat bioassay trial 25 rats were divided into 5 groups, each comprising of 5 rats in them.

Induction of Hypercholesterolemia

For induction of high cholesterol levels in rats, hypercholesterolemic diet was prepared and administered by using corn oil (10%), corn starch (64.5%), cholesterol (1.5%), protein (10%), mineral (3%), cellulose (10%) mineral and vitamin (1%).

Blood Sampling

The effect was authenticated by examining their total cholesterol content at interval of time 0 day, 15th day and 28th day. Rats were anesthetized by exposure to isoflurane and the blood samples were collected through cardiac puncture for biochemical analysis.
Biochemical Analysis

Serum lipid profile

The effect of selected defatted flax seed extract on the serum total cholesterol, low density lipoproteins (LDL), high density lipoproteins (HDL) and triglycerides levels of rats were determined by using method as described by (El-Newary et al.,2018).

Antioxidant status

The effect of defatted flax seed extract on the glutathione contents of rats was determined by following the protocol of Butler described by Sultan et al. Furthermore; the antioxidant enzyme assays include the estimation of concentration of superoxide dismutase (SOD) and catalase. The measurement of SOD activity is based on the method described by Sultan et al (Sultan et al., 2015).

Statistical analysis

Data was analyzed statistically using Statistix (9.0) software. Results were expressed as mean ± S.D. ANOVA (analysis of variance) for completely randomized design (CRD) was performed and to assess the significance (P ≤ 0.05) differences among means Tukey’s test was applied.

RESULTS

Total cholesterol (TC)

The statistical analysis reported a significant (p≤0.05) effect of time interval (0, 14, 28 days) and treatment on the cholesterol levels of hypercholesterolemic experimental rats. The results of the effect of MADFEs in different concentrations (150, 300 & 450mg/kg B.W.) on the levels of cholesterol are presented in figure 2. The results for levels of total cholesterol displayed peak (16.79%) percent reduction in experimental group F2, followed by F3 (13.3%) and F1 (11.83%). In comparison to the control F, 194.28±1.23 mg/dl, concentration of cholesterol in groups F2, F3 and F0 were observed to be 153.52±2.55 mg/dl, 141.62±2.33 mg/dl, 158.22±1.25 mg/dl respectively, on 28th day of administration. The highest reduction was observed in F2 group.

Table 1 Analysis of low-density lipoproteins (LDL) in experimental rats

<table>
<thead>
<tr>
<th>Parameter</th>
<th>Treatments</th>
<th>Interval of days</th>
</tr>
</thead>
<tbody>
<tr>
<td>LDL (mg/dl)</td>
<td>0 day</td>
<td>15th day</td>
</tr>
<tr>
<td>N0</td>
<td>37.22±1.87</td>
<td>38.67±1.21</td>
</tr>
<tr>
<td>F0</td>
<td>105.43±2.34</td>
<td>110.25±1.76</td>
</tr>
<tr>
<td>F1</td>
<td>110.14±1.65</td>
<td>104.63±1.44</td>
</tr>
<tr>
<td>F2</td>
<td>103.30±1.55</td>
<td>96.89±2.12</td>
</tr>
<tr>
<td>F3</td>
<td>108.19±2.65</td>
<td>107.45±1.23</td>
</tr>
</tbody>
</table>

Low density lipoproteins (LDL)

The results of the statistical analysis displayed significant (p≤0.05) effect of treatments (150, 300 & 450mg/kg B.W.) and time intervals (0, 14, 28 days) on the levels of LDL in experimental rats. The results regarding the effect of MADFEs on the LDL content is presented in table 1. The LDL content was reduced from 106.77±2.04 mg/dl at 0 day to 104.80±1.63 mg/dl at 14th day and 99.98±2.26 mg/dl at 28th day of administration, respectively. Moreover, the LDL concentration observed in F3 (110.77±2.36 mg/dl) was decreased to 104.16±1.81 mg/dl (F2), 96.80±1.99 mg/dl (F1), and 103.67±1.78 mg/dl (F0), respectively. The percent reduction in comparison to hypercholesterolemic control F0 was 12.68%, 11.2% and 11.8%; portraying highest reduction in F2.

Table 2 Analysis of high-density lipoproteins (HDL) in experimental rats

<table>
<thead>
<tr>
<th>Parameter</th>
<th>Treatments</th>
<th>Interval of days</th>
</tr>
</thead>
<tbody>
<tr>
<td>HDL (mg/dl)</td>
<td>0 day</td>
<td>15th day</td>
</tr>
<tr>
<td>N0</td>
<td>79.63±2.16</td>
<td>83.68±2.55</td>
</tr>
<tr>
<td>F0</td>
<td>107.45±1.23</td>
<td>105.42±1.55</td>
</tr>
<tr>
<td>F1</td>
<td>105.43±2.34</td>
<td>101.25±1.76</td>
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<tr>
<td>F2</td>
<td>103.30±1.55</td>
<td>96.89±2.12</td>
</tr>
<tr>
<td>F3</td>
<td>108.19±2.65</td>
<td>107.45±1.23</td>
</tr>
</tbody>
</table>

High density lipoproteins (HDL)

The statistical analysis demonstrated significant (p≤0.05) effect on the HDL level in the experimental rats. The effect of MADFEs on HDL content in experimental rats are presented in table 2. In comparison to the results of hypercholesterolemic control F0 (75.420±21.83mg/dl), an elevation identified in the values of F1 (79.630±2.16 mg/dl), F2 (83.680±2.55mg/dl) and F3 (78.497±1.60 mg/dl), respectively. However, the HDL levels were increased from 74.600±1.8 mg/dl (0 day) to 79.075±2.3mg/dl (14th day) and 84.245±1.8 mg/dl (28th day), respectively. The percent elevation displayed as compared to F0, were 12.09 %, 14.71 % and 11.21%; highest for F2..
The statistical analysis reported that time intervals and treatment have significant ($p<0.05$) effect on the levels of total triglycerides in hypercholesterolemic rats. The results of triglyceride levels in rats are displayed in figure 3. A considerable percent decrease of 12.25 %, 15.41%, 9.89 % was observed in experimental rats. Maximum reduction was observed in group F₁.

<table>
<thead>
<tr>
<th>Parameter</th>
<th>Treatments</th>
<th>Interval of days</th>
<th>0 day</th>
<th>15&lt;sup&gt;th&lt;/sup&gt; day</th>
<th>28&lt;sup&gt;th&lt;/sup&gt; day</th>
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<tbody>
<tr>
<td>Triglyceride (mg/dl)</td>
<td>N₀</td>
<td>0 day</td>
<td>76.74±1.26</td>
<td>80.37±1.23</td>
<td>84.72±1.54</td>
<td>80.52±1.34</td>
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<tr>
<td></td>
<td>F₀</td>
<td>180.58±2.21</td>
<td>189.60±1.43</td>
<td>200.50±2.98</td>
<td>190.23±2.20&lt;sup&gt;a&lt;/sup&gt;</td>
<td></td>
</tr>
<tr>
<td></td>
<td>F₁</td>
<td>185.22±1.29</td>
<td>179.32±2.65</td>
<td>162.53±2.76</td>
<td>175.71±2.23&lt;sup&gt;b&lt;/sup&gt;</td>
<td></td>
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<tr>
<td></td>
<td>F₂</td>
<td>178.02±2.34</td>
<td>169.03±1.55</td>
<td>150.57±1.98</td>
<td>164.87±1.91&lt;sup&gt;c&lt;/sup&gt;</td>
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<tr>
<td></td>
<td>F₃</td>
<td>181.84±2.07</td>
<td>174.87±2.27</td>
<td>163.85±1.24</td>
<td>173.52±1.82&lt;sup&gt;b&lt;/sup&gt;</td>
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<tr>
<td></td>
<td>Means</td>
<td>181.41±1.97&lt;sup&gt;c&lt;/sup&gt;</td>
<td>177.47±1.97</td>
<td>169.36±2.21</td>
<td>169.36±2.21</td>
<td></td>
</tr>
</tbody>
</table>

* N₀ normal group with no extract and high cholesterol diet, F₀ = hypercholesterolemic control group (high cholesterol diet), Treatment groups F₁, F₂, and F₃ (extract & high cholesterol diet)

Statistical analysis revealed a significant ($p<0.05$) effect of time interval and treatment on SOD enzyme levels in experimental rats. The results of SOD levels in rats are displayed in figure 3. The levels of SOD enzyme elevated from 20.59±2.30 (0 day) to 28.59±2.30 (28<sup>th</sup> day), respectively. While, in comparison to F₀ (22.17±2.45 IU/L), there was an elevation displayed in F₁ (24.28±1.82 IU/L), F₂ (24.88±2.30 IU/L) and F₃ (23.76±1.41 IU/L), respectively. The percentage increase showed by the experimental groups were 8.69%, 10.89% and 6.69%; highest recorded for F₃.

The peak elevation in the levels of SOD was noted for the group F₁.

**Figure 3** Analysis of superoxide dismutase (SOD)

* N₀ normal group with no extract and high cholesterol diet, F₀ = hypercholesterolemic control group (high cholesterol diet), Treatment groups F₁, F₂, and F₃ (high cholesterol diet + extract)

**Catalase (CAT)**

The statistical analysis demonstrated a significant ($p<0.05$) effect of the variables (treatments and time intervals) on the catalase enzyme level in experimental rats. The results for the levels of catalase are mentioned in table 4. The catalase levels elevated from 21.81±1.51 IU/L (0 day) to 23.19±1.72 (14<sup>th</sup> day) and 25.82±2.11 (28<sup>th</sup> day), respectively.
**DISCUSSION**

Hypercholesterolemia or dyslipidemia referred as abnormal blood lipid/fat levels. Hypercholesterolemia termed as high levels of cholesterol in the blood. Hypercholesterolemia is a risk factor for many diseases like heart diseases, diabetes, obesity, coronary heart diseases.so their levels must be normal in the body.a study reported that lignin secoisolariciresinolic diglisucide (SDG) in mice reduced the accumulation of fat in liver, and reduce the visceral fat accumulation in hyperlipidemia, hypercholesterolemia, hyperinsulinaemia and hypertrieneptaemia (Dyck, Heigenhauser and Bruce, 2006). Present study reported the same that defatted flaxseed extract lowers the cholesterol levels in hypercholesterolemic rats. A study was conducted on rabbits and 5 groups were made and 40mg/ml flaxseed lignin (defatted flaxseed) was given to them and blood samples were analyzed after treatment. the results showed significant decrease in lipid profile and inhibit lipid peroxidation. They reported the mean of cholesterol 143.8±13.9 in rabbits. Similarly, Prasad in 2005 reported that flax lignin reduced the significant reduction in cholesterol levels with a dose of 40mg/kg b.w when administered to 60 days (Dodin et al., 2006). A study was conducted by Zang and his fellows, according to that a dose of 300mg SDG was given to the group and it showed significant difference in in TC and LDL levels. They reported the reduction in range of 22.0 to 24.38 % in TC and LDL levels over a dose of 600mg/kg b.w. for many weeks. Many studies have been done on defatted flaxseed. It was noted previously that consumption of 40 g flaxseed/d with lowers amount of lignin (21 mg) for 12 mouths did not showed more effect on cholesterol level in post-menopausal women. There are many evidences from literature that reports the lipid lowering effects of defatted flax lignin. Faseh. In this study, the results showed that flaxseed lignin (defatted) showed a reduction of 81% in total cholesterol and 52% in triglyceride levels as compared to whole flaxseed. It is evident that dietary supplementation with partially defatted flaxseed reduced serum concentrations of total cholesterol, LDL cholesterol, and Apo lipoprotein B compared with the control treatment.

In a study it was reported (a systematic review) in which 9 clinical trials were done 6 out 9 showed significant decrease in total cholesterol and LDL levels when administered with flaxseed supplementation (Bassett et al., 2011). It has been stated in literature many times that lignin SDG from defatted flaxseed possess hypercholesterolemic and anti-atherogenic effects and SDG is the most likely component that have lipid lowering effects and reduce the plasma lipid profile. And reported by scientists in the findings that consumption of flaxseed daily in diet reduce the cholesterol (TC) levels by 6-11% and LDL by 9-18% normal patients while the levels of reduction in TC was 5-17% in hypercholesterolemic subjects and it was evident by their study that flaxseed lignin provide lipid lowering actions. This finding is close to our finding as in our study it reduced to 16%. And the levels of cholesterol dropped more with defatted flaxseed as compared to whole flaxseed (Truan et al., 2012).

Hyperlipidemic effects of flax lignin is due to multi mechanisms that have been reported in earlier studies, flax lignin is involved in increase or decrease at levels 7α-hydroxylase that converts cholesterol to bile. A study showed that SDG from flaxseed have an effective scavenging property of reactive oxygen species that inhibit lipid peroxidation through metal ions chelation (Mohammedi et al., 2006). It was reported in study and it is evident that supplementation with defatted flaxseed was observed to have lowered total cholesterol, LDL cholesterol concentrations, and Apo lipoprotein B in comparison to the control treatment. Our findings on defatted flaxseed extract had shown positive effect on elevated LDL levels and suggest that extract of defatted flaxseed lowers the levels of LDL in the body. In view of earlier studies on hyperlipidemic adults experimented with 50g/day of partially defatted flaxseed meal reduces the cholesterol 4.6% and LDL by 7.6% in 3 weeks. Similarly, Hallund studied the direct effect of flax lignin extract on plasma lipid profile, whereas a study reported that 500mg/day of flaxseed extract of lignin, reduced the plasma lipids in healthy women over a period of 6 weeks diet (Hallund et al., 2006) Similarly zhong and his fellows in 2008 reported that SDG of dose 300mg and 600mg/day in adults having high lipid profile reduce the LDL levels by 24% and total cholesterol by 22%.

Lignin content of flaxseed appears to be a contributing factor for hyperlipidemic effect of flaxseed. Van Niekirk and his fellows reported in their studies that decrease in the serum TC and LDL levels is due to SEG content in Partial lignin present in Partially defatted flaxseed. Flaxseed lignin Known to reduce the hypercholesterolemic parameters that include TC, TG and LDL. Several mechanisms have been proposed regarding how flax lignin reduce the cholesterol levels. Increase in excretion of bile acids by flaxseeds ingestion augments the removal of bad cholesterol (LDL) and amend the liver metabolism in a way that enhance the LDL cholesterol removed by hepatocytes, and also increases the LDL receptors (Zatta et al., 2002). The other striking finding shows that flax lignans almost standardized the lipid profiles parameters in liver of rabbit, and become almost normal levels of control groups. The widely held studies report that flaxseed modestly reduces TC (6%-11%) and LDL cholesterol (9%–16%) levels in normolipidemic and hypercholesterolemic patients i.e. TC (4.6%–17%), LDL cholesterol (3.9%-10%) (Sawant et al., 2015).

Conversely, results in a study reported that the increase in HDL levels of rabbits administered with treatment for 60 days, and these were consistent with previous literature that serum HDL levels tend to be increased when hypercholesterolemic rabbits treated for 12 days. However, some studies suggest that the defatted flaxseed showed no effect on serum HDL levels. results of another study is showing these values of percentage reduction as Total C (mmol/L) 5.5 ± 1.2, HDLC (mmol/L) 1.4 ± 2.8, LDL-C (mmol/L) 9.7 ± 1.8, Tricylglycerol (mmol/L) 10.2 ± 4.8 .20.D.E.C inatra et al.,2006 reported the reduction in lipid profile in flaxseed and lignin was 80.78–7.02, HDL 41.57 ± 6.96, TG 49.54 ±6.57, LDL 0.68 ± 0.14. Irregular lipid and lipoprotein metabolism plays an important part in hypertensive rats as well (Sawant et al., 2015). Lipid peroxidation, oxidative stress and changes in the lipid metabolism is the reason behind decreased levels of HDL and increase in level of other cholesterol. The other studies showed the blood samples were collected from 58 male rats. CCl4 as toxin. catalase, superoxide dismutase (SOD), and peroxidase was measured CCL4 induced rats having dose of 2.0 g kg b.w. 15.07 ± 8.0 day 75.07 ± 1.30 day 13.17 ± 1.5 and 24.12± 1.20 25.01±1.08 23.76±1.41

**Table 4 analysis of catalase (CAT)**

<table>
<thead>
<tr>
<th>Parameter</th>
<th>Treatments</th>
<th>Interval of days</th>
<th>Means</th>
</tr>
</thead>
<tbody>
<tr>
<td>Catalase (IU/L)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>N0</td>
<td>0 day</td>
<td>15th day</td>
<td>28th day</td>
</tr>
<tr>
<td>13.17±1.23</td>
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<td>21.99±2.87</td>
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<tr>
<td>20.15±1.98</td>
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<td>25.01±1.08</td>
<td>23.76±1.41</td>
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<tr>
<td>Means</td>
<td>21.9±1.51</td>
<td>23.19±1.72</td>
<td>25.75±2.11</td>
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</table>

*0 normal group with no extract and high cholesterol diet, F0= hypercholesterolemic control group (high cholesterol diet), Treatment groups F1, F2, and F3 (high cholesterol diet +extract)
peroxidase by 43.6%, 47.86%, and 52.0%, respectively. The present study on defatted flaxseed showed more potential than other studies. Researchers claimed the clear results regarding flaxseed constituents and its antioxidant activity. Flaxseed constituents as an antioxidant as seen by renewal of liver enzymes, which were different from normal to one due to harmfulness induced by toxin (CCl₄). Hence it was essential to investigate the oxidant/antioxidant status of the rats. SOD levels in the study of flaxseed extract on winster rats was reported to have mean value of 9.585±0.361 for extract (flaxseed concentrate) FLC (500mg/kg) in myocardial damage in doxorubicin-induced rats along with high cholesterol levels (Zanwar et al., 2013).

CONCLUSION

Present findings demonstrate that defatted flaxseed extract lowers the elevated serum lipids level, hypercholesterolemia, hypertension, and provides significant reduction in TC, LDL, and triglycerides (TG) levels in the experimental rats. Microwave assisted extraction is found to be beneficial technique for extraction as it is cheap, less time consuming and efficient. Defatted flaxseed extract presents a source of valuable minerals, fiber, phenolic and antioxidants. In vitro & in vivo studies on defatted flaxseed extract proved its efficacy and therapeutic potential against hypercholesterolemia. It is the richest source of dietary fiber and lignin, SDG is chief lignin found in defatted flaxseed. Flaxseed or defatted flaxseeds have great therapeutic potential reported in previous studies; against hypercholesterolemia and other diseases also i.e. diabetes, obesity, heart disease, cancer and hormonal issues. Therefore present study done with them. It possesses antioxidant activity and this effect may reduce the risk of other lifestyle related ailments like diabetes, atherosclerosis & hypertension.

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