HYPOCHOLESTEROLEMIC EFFECT OF MICROWAVE ASSISTED DEFATTED FLAXSEED EXTRACT IN EXPERIMENTAL RATS

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ABSTRACT

The research was aimed to determine the hypcholesterolemic 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 lower 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 hypcholesterolemic 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, enteroiol

INTRODUCTION

Hyperlipidemia is termed as elevated serum total cholesterol, triglycerides, LDL and VLDL levels, along with decrease in HDL levels that 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. anti-tumor 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 (Mall 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 enterodiol and enterolactone 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).

Figure 1 Graphical representation of experimental design. Microwave assisted extraction of defatted flaxseeds and administration of extracts in different doses to the study rats. After the administration analysis of the blood samples were taken to determine the lipid lowering effect of MAE of defatted flaxseed.

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 flaxseed 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 ($P \leq 0.05$) differences among means Tukey’s test was applied.

RESULTS

Total cholesterol (TC)

The statistical analysis reported a significant ($p \leq 0.05$) 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 F$_2$, followed by F$_1$ (13.3%) and F$_3$ (11.83%). In comparison to the control F$_0$, 194.28±1.23 mg/dl, concentration of cholesterol in groups F$_1$, F$_2$ and F$_3$ were observed to be 153.52±2.55 mg/dl, 141.62±2.33 mg/dl, 158.22±1.25 mg/dl respectively, on 28$^{th}$ day of administration. The highest reduction was observed in F$_2$ group.

Low density lipoproteins (LDL)

The results of the statistical analysis displayed significant ($p \leq 0.05$) 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±1.63mg/dL at 14$^{th}$ day and 99.98±2.26 mg/dl at 28$^{th}$ day of administration, respectively. Moreover, the LDL concentration observed in F$_2$ (110.77±2.36 mg/dl) was decreased to 104.16±1.81 mg/dl (F$_3$), 96.80±1.99 mg/dl (F$_2$), and 103.67±1.78 mg/dl (F$_1$), respectively. The percent reduction in comparison to hypercholesterolemic control F$_0$ was 12.68%, 11.2% and 11.8 %; portraying highest reduction in F$_2$.

| Parameter        | Treatments | Interval of days | Means       |
|------------------|------------|------------------|-------------|
|                  |            | 0 day           | 15$^{th}$ day | 28$^{th}$ day |
| LDL (mg/dl)      | N0         | 37.23±1.87      | 38.67±1.21  | 38.60±1.87  | 38.16±1.65  |
|                  | F0         | 105.43±2.34     | 110.25±1.76 | 116.64±2.98 | 110.77±2.36 |
|                  | F1         | 110.14±1.65     | 104.63±1.44 | 97.72±2.31  | 104.16±1.81 |
|                  | F2         | 103.30±1.55     | 96.89±2.12  | 90.20±2.31  | 96.80±1.99  |
|                  | F3         | 108.19±2.65     | 107.45±1.23 | 95.38±1.46  | 103.67±1.78 |
|                  |            | 106.77±2.04$^a$| 104.80±1.63$^a$ | 99.98±2.26$^a$ |

$^a$N0 normal group (no extract and high cholesterol diet), F0= hypercholesterolemic control group (high cholesterol diet), Treatment groups F1, F2, and F3 (high cholesterol diet +extract; 150 mg/kg b.w, 300 mg/kg b.w and 450 mg/kg b.w respectively)

High density lipoproteins (HDL)

The statistical analysis demonstrated significant ($p \leq 0.05$) 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 F$_0$ (75.420±21.83mg/dl), an elevation identified in the values of F$_1$ (79.630±2.16 mg/dl), F$_2$ (83.680±2.25mg/dl) and F$_3$ (78.497±1.60 mg/dl). However, the HDL levels were increased from 74.600±1.8 mg/dl (0 day) to 79.075±2.3mg/dL (14$^{th}$ day) and 84.245±1.8 mg/dl (28$^{th}$ day), respectively. The percent elevation displayed as compared to F$_0$ were 12.09 %, 14.71 % and 11.21%; highest for F$_2$.
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 mean values of the TG triglycerides are mentioned in table 3. Total triglyceride levels presented in F0 (190.23±2.20 mg/dl) decreased to 175.71±2.23 mg/dl (F1), 164.87±1.91 mg/dl (F2) and 173.52±1.82 mg/dl (F3), respectively. Furthermore, triglyceride levels reduced from 181.41±1.97mg/dl at 0 day to 177.47±1.97mg/dl at 14th day and 169.36±2.21 mg/dl at 28th day, respectively. In comparison to hypercholesterolemia control (F0) considerable percent decrease of 12.25 %, 15.41 %, 9.89 % was displayed in experimental rats. Maximum reduction was observed in group F2.

### Table 3 Analysis of triglyceride (TG) in experimental rats

| Parameter      | Treatments | Interval of days | 0 day                  | 15th day               | 28th day               | Means      |
|----------------|------------|------------------|------------------------|------------------------|------------------------|------------|
| Triglyceride (mg/dl) | N0         |                  | 76.74±1.26             | 80.37±1.23             | 84.47±1.54             | 80.52±1.34 |
|                | F0         |                  | 180.58±2.21            | 189.60±1.43            | 200.50±2.98            | 190.23±2.20 ^* |
|                | F1         |                  | 185.22±1.29            | 179.37±2.65            | 162.53±2.76            | 175.71±2.23 ^b |
|                | F2         |                  | 178.02±2.34            | 166.03±1.55            | 150.57±1.98            | 164.87±1.91 ^c |
|                | F3         |                  | 181.84±2.07            | 174.87±2.27            | 163.85±1.24            | 173.52±1.82 ^b |
| Means          |            |                  | 181.41±1.97 ^*         | 177.47±1.97 ^*         | 169.36±2.21 ^c         |            |

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

### Superoxide dismutase (SOD)

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 28.31±0.82IU/L (F0) to 32.36±0.66IU/L (F1), 40.09±0.99IU/L (F2) and 33.51±0.51IU/L (F3) on 28th day of administration of MADFESs, respectively. Furthermore, the maximum percent increase was displayed in group F1 (29.38%), followed by F3 (15.5%) and F2 (12.5%). The peak elevation in the levels of SOD was noted for group F3.

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

* ^*N0 normal group with no extract and high cholesterol diet, F0= hypercholesterol 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 (14th day) and 25.82±2.11 (28th day), respectively. While, in comparison to F2 (22.17±2.45 IU/L), there was an elevation displayed in F1 (24.28±1.82 IU/L), F3 (24.88±2.30 IU/L) and F2 (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.
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 secoisolariciresin diglucoside (SDG) in mice reduced the accumulation of fat in liver, and reduce the visceral fat accumulation in hyperlipidemia, hypercholesterolemia, hyperinsulimia and hyperptertensiona (Dyck, Heigenhauser and Bruce, 2006).

These reactive oxygen species have been shown to decrease in the serum TC and LDL levels is due to SEG content in Partial lignin and defatted flaxseed as compared to whole flaxseed (Deyck, Heigenhauser and Bruce, 2006).

Similarly, Prasad in 2005 reported that flax lignin and secoisolariciresin diglucoside in cholesterol levels with a dose of 40mg/kg b.w when administered to 60 days (Dodin et al., 2006).

In our study it was conducted by Zang and his fellows, to 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 for 6 weeks. Many studies have been done on defatted flaxseed. It was noted previously that consumption of 40 g flaxseed/d with lower amounts of lignin (21 mg) for 12 months did not show more effect on cholesterol level in post-menopausal women. There are many evidences from literature that reports the lipid lowering effects of flax lignin, as defatted flax lignin have been shown in literature many times that lignin secoisolariciresin diglucoside (SDG) in mice reduced the serum TC and LDL levels (HDL-cholesterolemia, hypolipidemic model reported the reduction in lipid profile LDL and TG (Mahdiesen et al., 2005), it was reported in study and 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.

Similarly, Amin et al who found that Rats fed high fat diet for 12 weeks showed a significant increase in the lipid profile LDL and TG, also it was also found that Plasma triglyceride was lowered by about 85–90% in high-fat fed rats (30%) for 8 weeks. all these evidence showed that high levels of triglycerides linked to many diseases including hypercholesterolemia (Stuglin and Parasad,2005),flaxseed meal reduce the cholesterol levels i.e. Total cholesterol, mmol/L 5.5 ± 0.5 and Triglyceride, mmol/L 4.09 ± 0.34. Similar addition of flaxseed extracts has also been linked with betterment of lipid serum profiles. Addition of 20% flaxseed in rat diet lowered total cholesterol level, triglyceride (TG) level, and low-density lipoprotein (LDL) cholesterol level by 21, 34, and 23%,respectively(Matusiewicz et al.,2015). results of another study is showing these values of percentage reduction as Total C (mmol/L) 5.5 ± 1.2, HDL-C (mmol/L) 1.4 ± 2.8, LDL-C (mmol/L) 9.7 ± 1.8, Triglycerglycerol (mmol/L) 10.2 ± 4.8. It was reported the reduction in lipid profile by flaxseed and presented means was total cholesterol 60.87± 7.02, HDL 41.57 ± 6.96, TG 49.54 ± 5.67, 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 lipids. Several studies have shown the reduction in lipid profile LDL and TG, also it was also found that Plasma triglyceride was lowered by about 85–90% in high-fat fed rats (30%) for 8 weeks. all these evidence showed that high levels of triglycerides linked to many diseases including hypercholesterolemia

Some antioxidant properties of flaxseed have been studied in vitro and in vivo studies, many researchers have found that flaxseed contain high level of antioxidants improve human health and inhibit cancer cells, improve cardiovascular diseases and metabolic disorders like diabetes (Zuk et al., 2011).

Natural antioxidants are present in the form of Phenolics, coumarins, Xanthons, lignans, Tannins, lycopene and beta carotene (Jeong et al., 2004). It is evident in leaves, seeds, and oil of the plant material. It has been reported previously that antioxidants have ability to inhibit human cancer cells, improve cardiovascular diseases and metabolic disorders like diabetes. Several studies have shown the reduction in lipid profile LDL and TG, also it was also found that Plasma triglyceride was lowered by about 85–90% in high-fat fed rats (30%) for 8 weeks. all these evidence showed that high levels of triglycerides linked to many diseases including hypercholesterolemia

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 individuals 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 (Trawan et al., 2012).

Hypolipidemic effects of flax lignin is due to multi mechanisms that have been reported in earlier studies, flax lignin is involved in increase of LDL, whereas flax lignin has been found that lignins 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 moderately 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 60 days. However, some studies suggest that 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, HDL-C (mmol/L) 1.4 ± 2.8, LDL-C (mmol/L) 9.7 ± 1.8, Triglycerglycerol (mmol/L) 10.2 ± 4.8. It was reported the reduction in lipid profile by flaxseed and presented means was total cholesterol 60.87± 7.02, HDL 41.57 ± 6.96, TG 49.54 ± 5.67, LDL 0.68 ± 0.14. Irregular lipid and lipoprotein metabolism plays an important part in hypertensive rats as well (Sawant et al., 2015).

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peroxidase by 34.6%, 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 (CC14). Hence it was essential to investigate the oxidant/antioxidant status of the rats. SOD levels in the study of defatted 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 (Zawar et al., 2013).

CONCLUSION

Present findings demonstrate that defatted flaxseed extract lowers the elevated serum lipids, glutathione peroxidase and total antioxidant capacity. Flaxseed 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 had 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 carried 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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