

## Hypocholesterolemic effects of cold and hot-pressed linseed oil in a Wistar Rat Model

**Bandar H. Aloufi**

Department of Biology, Faculty of Science, Hail University, Saudi Arabia

E-mail : bandaraloufi@yahoo.com

Contact No. : +.....

Submitted : 10.07.2017

Accepted : 27.11.2017

Published : 30.12.2017

### Abstract

Hypercholesterolemia is metabolic deregulation of cholesterol levels in the blood and the major risk factor that precipitate coronary heart disease and atherosclerosis. This study aimed to compare the efficiency of cold and hot-pressed linseed oil to decrease the levels of total cholesterol and proteins in Wistar rats (n =60) weighing (225-252 g). Linseed oil was fed for seven weeks to Wistar rats and control. Many biochemical and physiological parameters were measured by automatic analyzer and the results were statistically analyzed by SPSS.12. In comparison to control rats showed highly significant decrease in values of following physiological parameters; triglycerides, total cholesterol, low density lipoprotein cholesterol, total protein. In contrast, marked increasing in the value of high density lipoprotein cholesterol. Highly significant decrease in rats fed with cold pressed linseed oil blood measurements of triglycerides, total cholesterol, and low density lipoprotein cholesterol. In contrast, marked increasing in the value of high density lipoprotein cholesterol when compared with rats fed with hot pressed linseed oil. These findings indicate that diets containing Linseed oil significantly improved the physiological parameters of rats. We suggest that Linseed oil as part of food might improve blood parameters and increase high density lipoprotein cholesterol in rats. We further suggest that Linseed oil supplementation act as antioxidant agents, and an excellent adjuvant therapy for rats.

Key words : Linseed oil, on lipid Metabolism, cholesterol.

### INTRODUCTION

Flax (common flax or linseed) or, belonged to family Linaceae. The Genus comprised of about 200 species<sup>[1]</sup>. Linseed contains 35-45% oil, 28% soluble dietary fiber, and 21% protein. Linseed comprises nutritional valuable components such as protein (200-240 g/kg), dietary fiber (250-280 g/kg) and flax oil (350-450 g/kg). The health benefits are related with the ingestion of polyunsaturated fatty acids (PUFA) and dietary fiber<sup>[1-3]</sup>.

Linseeds among unique oil seed crops because of its exceptionally high content of  $\alpha$ -linolenic acid (45 to 52% of its oil) (ALA), each tablespoon of ground linseed contains about 1.8 grams of plant omega- (E1)<sup>[4-6]</sup>. High amounts of ALA, were derived from Soybean.

Omega-3 fatty acid plays an important role in prevention or treatment of cardiovascular disease, hypertension, atherosclerosis, cancer neurological disorders and inflammatory disease. More omega-3 fatty acids intake decreases serum cholesterol which beneficially affects blood pressure, skin diseases, thrombosis atherosclerosis and diabetes, arterial compliance and hyperlipidemia response<sup>[7-8]</sup>. Linseed- but not linseed oil - contains soluble fiber. It might cause diarrhea, cramping, wind, and bloating. Large amounts of flaxseed, especially when not taken with enough water, can cause constipation and even bowel obstruction, flatulence, stomach pains, nausea and constipation<sup>[9-13]</sup>.

This study aimed to compare the efficiency of cold pressed Linseed oil and hot pressed linseed oil to decrease the levels of total cholesterol, hypocholesterolemic Effects and total lipids in a Wistar rats model.

### MATERIALS AND METHODS

#### MATERIALS

##### Linseed Oil:

Cold pressed Linseed Oil was obtained by cold pressing method which means the oil is pressed by great pressure of physical machinery under low temperature, so it is called cold pressing method. Cold pressed Linseed Oil was obtained by from Bio Oils, Canterbury, New Zealand, Containing the Essential Fatty Acids Omega 3, 6 and 9; linseed Oil is nature's richest source of Omega 3. Linseed oil was obtained from non-genetically modified seed.

Hot pressed flaxseed oil was commercially purchased from local market. The oil was extracted by hot pressing method. The oil is produced by physical pressing from oil crops after high temperature frying or steaming. It is the traditional pressing technology with high yield efficiency.

##### Animals

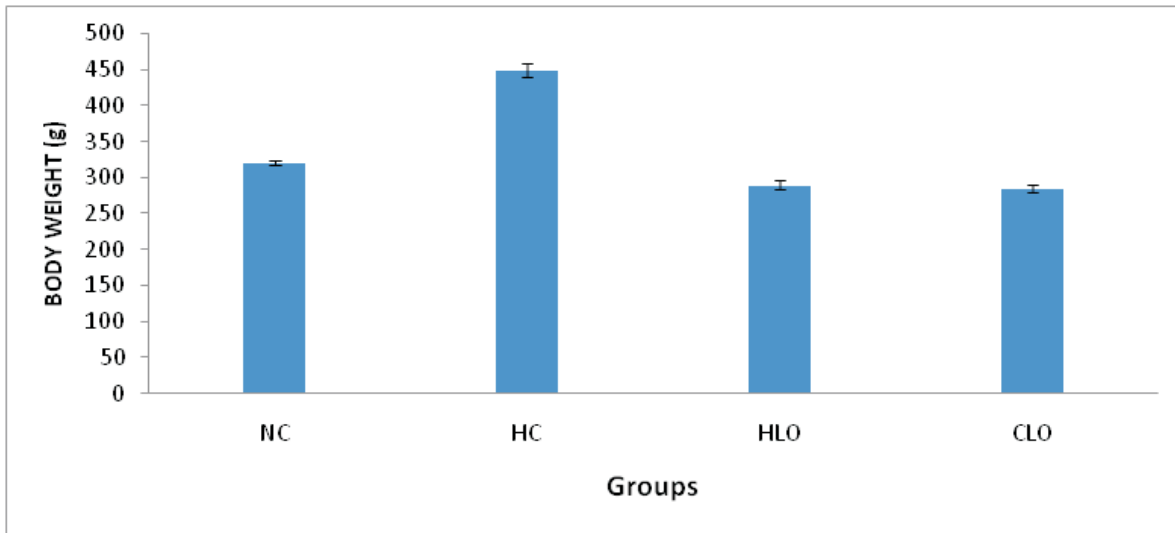
Healthy young adult male Wistar rats weighing (225-252 g) were obtained from The Animal physiology Lab of Faculty of Science Hail University.

The rats were housed in well-aerated individual cages and maintained in a temperature-controlled room (24 ± 1 °C) with a 12 h light/12 h dark cycle, 55 ± 10 % humidity. They were fed with normal commercial chow and water ad libitum. Throughout the experiments, animals were processed according to the suggested international ethical guidelines for the care of laboratory animals.

#### METHODS

##### Experimental design:

A total of 60 rats were used in the experiment. The rats were divided into 4 groups of 15 animals each as follows:



**Figure 1:** Effects of linseed oil supplementation on body weight gain, after 7 weeks of treatment.

Group 1: Normal control (normal rats) received normal commercial chow and water ad libitum.

Group 2: fed hypercholesterolemic diet (standard diet + 2% cholesterol) and water ad libitum.

Group 3: hot pressed linseedoil Group received diet was enriched with 5g/100g diet of hot pressed linseedoil.

Group 4: cold pressed linseedoil Group received diet was enriched with 5g/100g diet of cold pressed linseedoil.

#### Blood collection and determination of physiological parameters

At the end of experimental period, blood samples were collected from retro-orbital eye plexus [3].

Each sample was collected into both heparinized tubes to obtain the plasma and into a dry clean centrifuge glass tube without any coagulation to prepare serum. Blood was left for 15 min at room temperature, then the tubes were centrifuged for 15 min at 3000 rpm and the clean supernatant serum was kept frozen at -20 °C until the time of analysis for different biochemical analyses, prior immediate determination of triglycerides, cholesterol, high density lipoprotein HDL-cholesterol (HDL-C), low density lipoprotein LDL-cholesterol (LDL-C).

All of these parameters were measured using an automatic analyzer (Architect c8000 Clinical Chemistry System, USA).

#### Statistical analysis

Statistical analyses were performed using SPSS package for Windows version 13.0. Data are expressed as mean  $\pm$  SE. One-way ANOVA and two-way ANOVA were used to analyze differences among groups. Post-hoc analyses of significance were made using least-significant difference (LSD) test. Differences between groups were considered statistically significant at  $p < 0.05$ .

## RESULT

### Blood glucose

The mean values of blood glucose of control and treated experimental groups are presented in Table 1. No significant differences were observed in blood glucose level of normal rats

fed on diets containing the oil of linseed when compared with those rats fed on the control diet after 7 weeks of treatment.

### Blood triglyceride, cholesterol, LDL-C and HDL-C

The changes in the levels of serum lipids in control and experimental groups are illustrated in Table 1.

The rats were treated with linseed oil resulted in a significant ( $p < 0.01$ ) decrease in the levels of triglycerides, cholesterol and LDL-cholesterol compared to untreated rats. While HDL-cholesterol level was significantly ( $p < 0.01$ ) increased.

The rats exposed to the diets containing the linseed oil for 7 weeks had higher blood HDL-cholesterol than those of the control group ( $p < 0.05$ ).

We have found in this study that highly significant decrease in the group fed with cold pressed linseed oil in the measurements of triglycerides, total cholesterol, and low density lipoprotein cholesterol. In contrast, marked increasing in the value of high density lipoprotein cholesterol when compared with rats which fed with hot pressed linseed oil.

## DISCUSSION

Several studies demonstrated that a variety of herbal extracts effectively lowered the glucose level in normal experimental animal and STZ-induced diabetes mellitus rats.<sup>[3,6,8]</sup>

In the present study, many blood physiological parameters have been improved in rats fed on diets containing the oil of linseed when compared with those rats fed on the control diet after 7 weeks of treatment.

On the other hand, several researchers also concluded that *Linum usitatissimum* linseed oil significantly lowered the blood sugar level in glucose-fed and adrenaline induced hyperglycemic rats<sup>[8,14]</sup>.

However, many works observed that the association of hyperglycaemia with an alteration of lipid parameters presents is a major risk for cardiovascular complications in diabetes. Many secondary plant metabolites have been reported to possess lipid-lowering properties<sup>[15-18]</sup>.

**Table 1:** Effects of linseed oil supplementation on blood glucose, triglyceride and total lipid After 7 weeks of treatment.

Treatments	Glucose (mg/dl)	Plasma TG (mg/dl)	Total lipid (mg/dl)
NC	97.18±9.50	64.10±3.46	259.63±2.51
HC	108.50±3.74*	148.21±11.62*	302.11±3.80*
HLO	95.13±2.45#	61.90±2.15*#	230.96±7.01*#
CLO	90.77±3.92#	56.81±3.47*#	217.28±5.97*

The number of animals was 10 for each group

NC: Normal control, HC: hypercholesterolemic diet control, HLO: Hot-pressed linseed oil & CFO: cold-pressed linseed oil,

All values are expressed as means ± SE.

Significantly different from normal control (\* p < 0.05).

Significantly different from: hypercholesterolemic diet control (# p < 0.05).

**Table 3:** Effects of linseed oil supplementation on Cholesterol, HDL-C and LDL-C after 7 weeks of treatment.

Treatments	Cholesterol (mg/dl)	HDL-C (mg/dl)	LDL-C (mg/dl)
NC	88.38±3.31	39.54±2.28	38.52±3.25
HC	253.45±6.49*	21.84±3.92*	90.57±8.19*
HLO	62.21±1.25*#	43.62±2.75*#	31.51±2.67*#
CLO	51.01±9.77*#	50.62±5.53*#	29.44±2.37*#

The number of animals was 10 for each group

NC: Normal control, HC: hypercholesterolemic diet control, HLO: Hot-pressed linseed oil & CFO: cold-pressed linseed oil,

All values are expressed as means ± SE.

Significantly different from normal control (# p < 0.05, ## p < 0.01 and ### p < 0.001).

Significantly different from: hypercholesterolemic diet control (# p < 0.05).

The serumcholesterol and triglycerides were significantly decreased in diabetic rats supplemented with of linseedoil. The oilsupplementation also result the significant attenuation in the levels of HDL-cholesterol and LDL-cholesterol in serum toward the control level which again strengthen the hypolipidaemic influence of these oils. A variety of derangements in metabolic and regulatory mechanisms, due to insulin deficiency, is responsible for the observed accumulation of lipids<sup>[19-20]</sup>.

The impairment of insulin secretion results in enhanced metabolism of lipids from the adipose tissue to the plasma. Further, it has been reported that diabetic rats treated with insulin

show normalized lipid levels<sup>[3,9,21]</sup>.

We suggest that the present effects of these oils-treated diabetic rats may be due to its role in normalization of insulin secretion, lowering activity of lipid biosynthesis enzymes, especially cholesterol and or lowering level of lipolysis.

Moreover, many minor components of foods, such as secondary plant metabolites, have been shown to alter biological processes which may reduce the risk of chronic diseases in humans. popularly known as linseeds an indigenous plant widely available in India and Burma. Different parts of this plant have been reported to have antiseptic, wound healing and skin disease curing activity<sup>[22-24]</sup>.

Several studies demonstrated that water soluble portion of alcoholic extract of leaves of *Linum usitatissimum* possesses significant anti-inflammatory, antiserotonin, antifertility and hepatoprotective activity<sup>[10-13]</sup>. Significant hypolipidemic activity in rats fed on atherogenic diet and antihyperglycemic as well as hypotensive activity have also been reported by us<sup>[8]</sup>.

In the current research illustrated that diets containing linseed oil significantly improved the physiological parameters of rats and hypolipidemic and antioxidant activities of cold and hot-pressed linseed oil in high-cholesterol fed rats. Linseed and linseed oil has also been reported to act as anti-arrhythmic, anti-atherogenic and anti-inflammatory agent in addition to its improving of cardiovascular functions<sup>[6-9]</sup>.

However, many minor components of foods, such as secondary plant metabolites, have been shown to alter biological processes which may reduce the risk of chronic diseases in humans. *Linum usitatissimum* popularly known as linseed is an indigenous plant widely available in India and Burma. Different parts of this plant have been reported to have antiseptic, wound healing and skin disease curing activity<sup>[22-24]</sup>.

We have investigated in this study that highly significant decrease in rats fed with cold pressed linseed oil blood measurements of triglycerides, total cholesterol, and low density lipoprotein cholesterol. In contrast, marked increasing in the value of high density lipoprotein cholesterol when compared with rats fed with hot pressed linseed oil and these results indicated that the process method has a significant effect on the aroma quality of FSO and may be helpful in evaluating aroma quality as Chang Q. Wei et al reported in 2015<sup>[25]</sup>.

## CONCLUSION

The present data suggest that using linseed oil improve blood parameters. The responses in blood parameters in these animals are also demonstrated that *Linum usitatissimum* oil supplementation may act as antioxidant agents and could be an excellent adjuvant support in the therapy of hypercholesterolemia and hyperlipidemia.

## REFERENCES

1. Biswas K, Chattopadhyay I, Banerjee R, Bandyopadhyay U. Biological activities and medicinal properties of (*Azadirachta indica*). *Cur. Sci.*, 2002; 82(11): 1336-1345.
2. Chattopadhyay RR, Chattopadhyay RN, Nandy AK, Podder G, Maitra SK (1993). Preliminary report on antihyperglycemic effect of a fraction of fresh leaves of *Azadirachta indica* (Beng. Flaxseed). *Bull. Calc. Sch. Trop. Med.* 1993; 35: 29-35.
3. Eldemerdash FM, Yousef M I, Abou El Naga NI. Biochemical study on the hypoglycemic effects of onion and garlic in alloxan induced diabetic rats. *Food Chem. Toxicol.* 2005; 43: 57-63.
4. Srivastava V, Varma V, Raval S. Antidiabetic and adaptogenic properties of *Mamodica charantia* extract: An experimental and clinical evaluation. *Phytother. Res.* 1993; 7: 285-289.
5. "Essential Fatty Acids". & Micronutrient Information Center, Oregon State University, Corvallis, OR. May 2014. Retrieved 24 May 2017.
6. Hussein HK and Abu-Zinadah OA. Antioxidant effect of curcumin extracts in induced diabetic Wistar rats. *Int. J. Zool. Res.* 2010; 6(4): 266-276
7. MacLean CH, Newberry SJ, Mojica WA, Khanna P, Issa AM, Suttorp MJ, Lim YW, Traina SB, Hilton L, Garland R, Morton SC. "Effects of omega-3 fatty acids on cancer risk: a systematic review." *JAMA: The Journal of the American Medical Association.* 2006; 295 (4): 403-15.
8. Chattopadhyay RR, Chattopadhyay RN, Nandy AK, Podder G, Maitra SK. Preliminary report on antihyperglycemic effect of a fraction of fresh leaves of *Azadirachta indica* (Beng. Flaxseed). *Bull. Calc. Sch. Trop. Med.* 1993; 35: 29-35.
9. Massing MW, Sueta CA, Chowdhury M, Biggs DP, Simpson RJ. Lipid management among coronary artery disease patients in diabetes mellitus or advanced age. *Amer. J. Cardiol.* 2001; 87: 646-664.
10. Rajasekaran SK, Avi K, Sivaganam K, Subramanian S. Beneficial effects of aloe vera leaf gel extract on lipid profile status in rats with streptozotocin diabetes. *Clin. Exp. Pharmacol. Physiol.* 2006; 33: 232-237.
11. Pathak RM, Ansari S, Mahmood A (1981). Changes in chemical composition of intestinal brush border membrane in alloxan induced chronic diabetes. *Indian J. Exp. Biol.*, 19: 503-505.
12. Rajasekaran S, Sivaganam K, Subramanian S. Antioxidant effect of Aloe vera gel extract in streptozotocin-induced diabetes in rats. *Pharmac. Report.* 2005; 57: 90-96.
13. Ravi K, Ramachandra B, Subramanian S. Protective effect of *Eugenia jambolana* seed kernel on tissue antioxidants in streptozotocin induced diabetic rats. *Biolog. Pharm. Bull.* 2004; 27: 1212-1217.
14. Sekar DS, Sivaganam K, Subramanian S. "Antidiabetic activity of *Momordica charantia* seeds on streptozotocin induced diabetic rats, *Pharma.* 2005; 60(5): 383-387.
15. Sharma SB, Nasir A, Prabhu KM, Murthy PS, Dev G (2003). Hypoglycaemic and hypolipidemic effect of ethanolic extract of seeds of *Eugenia jambolana* in alloxan-induced diabetic rabbits. *J. Ethnopharmacol.*, 85: 201-206.
16. Srivastava V, Varma V, Raval S. Antidiabetic and adaptogenic properties of *Mamodica charantia* extract: An experimental and clinical evaluation. *Phytother. Res.* 1993; 7: 285-289.
17. Yehuda, Shlomo; Rabinovitz, Sharon; Mostofsky, David I. (2005). "Mixture of essential fatty acids lowers test anxiety". *Nutritional Neuroscience.* 8 (4): 2657. One. 7 (9): e46028. ISSN 1932-6203.
18. Chang Q. Wei, Wen Y. Liu, Wan P. Xi, Dong Cao, Hui J. Zhang, Ming Ding, Lu Chen, Ya Y. Xu and Ke X. Huang. Comparison of volatile compounds of hot-pressed, cold-pressed and solvent-extracted flaxseed oils analyzed by SPME-GC/MS combined with electronic nose: Major volatiles can be used as markers to distinguish differently processed oils. *Eur. J. Lipid Sci. Technol.* 2015; 117, 320330.
19. Xin, Wei; Wei, Wei; Li, Xiaoying. "Effect of fish oil supplementation on fasting vascular endothelial function in humans: a meta-analysis of randomized controlled trials". *PLoS One.* 2012; 7(9): e46028. ISSN 1932-6203

20. Yoshida M, Kimura H, Kyuki K, Ito M. Effect of combined vitamin E and insulin administration on renal damage in diabetic rats fed a high cholesterol diet. *Biol. Pharm. Bull.* 2005; 28: 2080-2086.
21. Zimmer, Carl. "Inuit Study Adds Twist to Omega-3 Fatty Acids' Health Story". *New York Times*. Retrieved 11 October 2015.
22. Su, Kuan-Pin; Huang, Shih-Yi; Chiu, Chih-Chiang; Shen, Winston W. "Omega-3 fatty acids in major depressive disorder". *European Neuropsychopharmacology*. 2003; 13 (4): 26771.
23. Van Dam RM, Rimm EB, Willett WC, Stampfer MJ, Hu FB (2002). Dietary patterns and risk for type 2 diabetes mellitus in U.S. men. *Ann. Int. Med.*, 136:201-209.
24. Wang L, Folsom AR, Zheng ZJ, Pankow JS, Eckfeldt JH . Plasma fatty acid composition and incidence of diabetes in middle-aged adults: the Atherosclerosis Risk in Communities (ARIC) Study. *Am. J. Clin. Nutr.* 2003; 78:91-98.
25. Chang Q, Wei, Wen Y, Liu, Wan P, Xi, Dong Cao, Hui J, Zhang, Ming Ding, Lu Chen, YaY. Xu and Ke X. Huang (2015). Comparison of volatile compounds of hot-pressed, cold-pressed and solvent-extracted flaxseed oils analyzed by SPME-GC/MS combined with electronic nose: Major volatiles can be used as markers to distinguish differently processed oils. *Eur. J. Lipid Sci. Technol.* 2015; 117, 320330.