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Body Weight, Hematological and Biochemical Parameters of Wistar Albino Rats Fed with Chia Seed Concentrate

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Authors' contributions

This work was carried out in collaboration among all authors. All authors read and approved the final manuscript.

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ABSTRACT

Chia seeds are among commonly used natural remedies to control obesity, diabetes mellitus and cardiovascular diseases. A research was conducted to evaluate the effects of chia seeds on Body weights, hematological parameters, blood glucose, and lipid profiles in Wistar rats. In the current study, Wistar rats were fed on chia seed powder mixed with broiler chicken feed mash at concentrations of 0, 5, 10, 15 and 20%. Body weights, hematological parameters, blood glucose, and lipid profiles were monitored for 28 days. Results indicate that for the treated groups, there was significant increase in white blood cell (WBC) count and lymphocytes in a dose dependent manner ($R^2 = 0.87$; p = 0.032). Plasma glucose and lipid profiles also decreased significantly in an dose dependent manner, whereby, for glucose, group 4 rats-decreased from 138.5 ± 0.8 to 80.3 ± 12.9

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mg/dL from day 0 to day 28 of treatment. The results show no significant difference on weight gain in chia treated groups compared to control group. There were no significant variations in the other hematological parameters. In conclusion, findings in this study illustrated that chia seeds can be the good candidate for controlling blood glucose, lipid profiles and improving hematological parameters and also body weight in animals.

Keywords: Chia seeds; blood glucose; lipid profile; rats.

1. INTRODUCTION

Recent reports from the WHO [1] have indicated increased mortalities of up 17.9 million deaths per year attributable to metabolic disease conditions such as obesity, diabetes mellitus and cardiovascular diseases, where, about 1.5 millions of the deaths were caused by diabetes mellitus. Treatment of metabolic disorders often requires changes of lifestyle habits such as consumption of healthy diets and engaging in routine physical exercises [2,3]. However, it can be challenging for patients to adopt this lifestyle and faithfully adhere to it.

Despite the several industrial medications available to treat some metabolic disorders, there are still many challeges in treatment and management of these conditions [4,2]. For instance, finding the right medication and dosage for each patient can be a challenge. Additionally, some patients have adverse reactions to certain medications while others have limited access to healthcare resources, [5,6]. These challenges necessitate for search of alternative therapeutic functional approaches including foods supplementation towards preventing and controlling these conditions and other associated health risks. Several herbs, spices and natural remedies have been used as alternative therapeutics to these medications [7,8,9]. Chia seeds are among the commonly used natural remedies for controlling obesity, diabetes mellitus and cardiovascular diseases [10,11,12]. The seeds have been in used for instance for medicinal purposes by Mexican communities, among others since 3500 BC [13]. Nowadays, Chia seeds have extended use beyond medicine as used by ancient communities and have many industrial uses including; Oil industry to produce edible oil and mayonnaise, beverage industry where the seeds and products are used to enrich beverages, milk industry where it is use in butter, cheeses, and fresh dairy products such as yoghurt, packaging industry where the seeds are used to produce biodegradable films, meat, snacks and extrusion industries among others [14,15,16,17].

is an annual flowering plant that Chia belongs to a family Lamiaceae. The seeds are known to contain mainly; protein, fat. carbohydrate, dietary fibre, ash and dry matter contents [18,19]. They also contain phenolic compounds. minerals and omega-3 fattv acids, and *a*-linolenic acid [20]. Omega-3 fatty acid are essential fatty acids known to have a large number of physiological functions in body [21,22], human for instance. they attributable function as antioxidants to chlorogenic acid, caffeic acid. mvricetin. quercetin, and kaempferol [23]. These agents are believed to have renal, cardiac, hepatic effects. dvslipidemia. protective anti anti hypertension, anti-ageing and anti-carcinogenic characteristics [24,10].

Chia seed are also known to improve insulin sensitivity, maintenance of bone mass. inflammation of the tissue decrease and decrease vascular reactivity [25,26,7]. They also contain high fiber content important in the control body weight gain and obesity [27,28]. Chia seeds however have a hardy seed coat which makes the hydrolysy of the coat lipids and proteins difficult hence difficult to release the bioactive components of the seed matrix. This makes an overall difficulty in bioavailaility of nutrients from the seeds [29]. In addition, they are small in size and lowly cohesive, making them difficult to incoropearate into food and feed products [30]. Processing of the seeds could simplify the release of the valuable chia seeds nutrients making them bioavailable. Grinding of the seeds is one of the simple ways to expose the seeds matrix from the hard seed coat to make the nutrients bioavailable [30] while preserving them. In this research we used chia seed powder after sun drying and mixing it with commercial broiler chicken mash to compose experimental animal feeds.

In this study, we evaluated the effect of ground dried chia seed on body weight, selected hematological and biochemical parameters using wistar albino rats as model animals.

2. MATERIALS AND METHODS

2.1 Study Area

The study was conducted at College of Veterinary Medicine and Biomedical Science (CVMBS), Sokoine University of Agriculture (SUA), Morogoro.

2.2 Research Design

An experimental study was done where, mature healthy male and female wistar albino rats were fed on different percentages of chia seed meal and broiler chicken mash as basal diet to study the effects of the seeds on body weight and selected hematological parameters.

2.3 Chia Seed Meal Preparation

Five chia seed feed formulations were prepared for the experiments at concentrations of 0, 5, 10, 15 and 20% in commercial broiler chicken feeds mash. The above formulations were prepared by addition of 0, 50, 100, 150, and 200g of chia seed powder to make 1,000g of the commercial chicken mash different broiler for feed formulations making up five formulations refered to as formulation 0 (F0) to formulation 5 (F4). The feeds were compounded three times (once a week) and stored in cool dry place without direct sun light throught the experiment period.

2.4 Preparation of Experimental Animals

A total of 100 mature male and female wistar albino rats weighing between 180 to 200g were used for the experiments. The rats were divided into five subgroups each with 20 wistar albino rats, designated as group 0 (G0) to group 4 (G4). G0 was used as the control group while G1-G4 were the treatment groups. All the rats were housed in zoo-sanitary cages and had full access to potable tap drinking water and the formulated broiler chicken mash feeds with chia seed powder. Feeding the rats was as follows: G0 (Control group) was the control group and was fed on F0. The rest of the groups were fed on F1 for G1 to F4 for G4 respectively daily for 28 days. The room temperature and humidity conditions in the experimental rooms were adopted to the general environmental weather conditions at averages of 26°C temperature and 75% humidity of the the Sokoine University of Agriculture environmental conditions. Dark and light conditions also adopted the environmental which approximated to 12 hours of light (day) and 12

hours of darkness (night). Windows to the experimental house were open and closed to provide the light and dark cycles.

2.5 Data and Samples Collection

Data was collected on feed intake and body weights. These were recorded daily for all the rats in all cages during the entire period of the experiment. Whole blood samples were also collected.

For the weights, the rats were weighed to the nearest grams using the Mettler PE 6000® Switzerland, weighing scale. For the blood samples, approximately 4 ml of intracardial blood samples were collected using procedures explained by Parasuraman et al. [31] an approach to collection of blood samples in small laboratory animals. The blood samples were collected on days 0, 7, 14, 21 and 28 of the experiment. Briefly, the rats were anasthetised in chloroform where a piece of gauze was soaked in 1 ml of chloroform and introduced into a bell jar and closed with a lid. Each time of sampling, a rat was introduced into the jar for about sixty seconds to anasthesize the rats. Blood samples were collected through cardiac puncture of three rats from each group on every sampling day. About 4ml of blood was collected from each rat through cardiac puncture where about 1ml of the blood were transferred into vacutainer tubes containing ethyl diamine tetraacetate (EDTA) for haematological parameters analyses. 2ml of the blood samples from the respective rats and groups were immediately centrifuged at 3,000 revolution per minute (rpm) for 10 minutes to obtain fresh plasma. The plasma samples were used for analyses of hematological and parameters biochemical usina spectrophotometer (Cole Patner 1,100 rs). All sample collections were done in the mornings between 7 and 9 AM of the sampling day in order to reduce on diurnal variations that could affect the haematological parameters. Blood samples were collected in vacutainer tubes with anticoagulants later and analysed for haematological parameters using the MS[™]4-s, (MS laboratories, France) automated hematological analyzer. For glucose assays, the blood samples were collected in sodium florinated (NaF) vacutainer tubes to preserve glucose while for the other biochemical parameters, the blood samples were collected in heparinized vaccutainer tubes. The blood samples were centrifuged to obtain plasma which were stored at 4°C prior to assays.

2.6 Determination of Plasma Glucose and Lipid Profiles

Blood glucose and plasma total lipid profiles were done by assays of alucose and lipids using the Trinders methods [32] in the blood plasma. Briefly. for glucose assavs. the Erba® Mannheim,(Erba Lachema s.r.o., Karásek 1d, 621 00 Brno, CZ) glucose kit which employs the glucose-oxidase principle were used. Plasma samples and Trinders reagents were added together and mixed thoroughly for about five seconds in a vortex mixer and immediately incubated for five minutes at in dark at 37°C. Together with glucose standard, the mixtures were immediatetly read for absorbances in the **ColePalmer**® 1100RS Spectrophotometer (Product of the United Instrument Inc.).spectrophotometer following blanking of the spectrophotometer with distild water. Glucose concentrations were determined using the formula. "Concentration of glucose = Absorbance Sample/Absorbance of of Standard Concentration of Standard". For the lipid profiles, the Trinders calorimetric enzymatic methods by (CELLBIOLABS INC., Creating Solutions to Lifescience Research) kits for HDL and LDL/VLDL cholesterol were used. Briefly, samples and cholesterol standards were incubated for 45 minutes and then read in the calorimeter. Concentration of the lipids in the samples were compared to the known stardard cholesterol curve. Values were expressed on mg/dl.

2.7 Data Analysis

Data were statistically analyzed to check for differences of the means among the groups using one way analysis of variance (ANOVA). Differences were considered significant when *P values* were equal to or less than 0.05. Differences between the means of the groups were also analyzed for significancy using the student's *t*-test and a Tukey's Honesty Significance Difference (Tukey HSD) where differences were considered significant at P equal or less than 0.05 .The means were expressed with there corresponding standard error of the means (mean \pm SE). The data were analyzed in the statistical package for social sciences (SPSS) version 29, 2022 [33].

3. RESULTS

3.1 Body Weight

Feedd consumption and body weight measurements were carefully monitored in all groups of rats throughout teh experimental period. Feeds consumption and body weight gain results for the control and experimental groups are indicated in Table 1. From the results, mean body weights of the rats in the control group were slightly lower than, but not significantly different (p=1.069) from the treatment groups.

3.2 Hematological Results

Generally there were no signicant differences (P>0.05) in hematological parameters among the rats groups as inidcated in Table 2. the for red blood cells (RBC) count, blood hemoglobin concentration (Hb) and hematocrit (Hct). But there was significant dose dependent increase (p<0.05) in the white blood cell count (WBC), lymphocytes (LYM) and monocytes (MON) concentrations. The mean platelet volume (MPV) and platelets (PLT) were observed to be significantly lower than (p = 0.008) the control group G0.

Plasma glucose concentration: At baseline, the plasma glucose levels were not significantly different between the groups. The level of plasma glucose was observed to decrease significantly (p<0.05) in all treated groups in a dose dependant manner. The decrease was more prominent on 7th day of G3 and G4 (R²=0.78 p = 0.032).

Table 1. Means ± SE of body weights (g) for control of the rats which were fed on basal diet and treatment groups which were fed on chia seed feed formulations for 28 days

Groups	Treatment time in days						
	0	7	14	21	28		
G0	155.9 ± 16.5	193.8 ± 14.5	177.9 ± 12.3	179.7 ± 12.6	185.3 ± 12.3		
G1	190.6 ± 15.2	182.5 ± 17.7	180.5 ± 12.1	173.1 ± 10.5	175.1 ± 14.2		
G2	178.7 ± 1 4.3	183.4 ± 12.8	170.3 ± 17.5	175.0 ± 14.4	167.5 ± 21.7		
G3	189.9 ± 12.9	183.7 ± 17.8	184.0 ± 12.3	178.6 ± 21.2	165.9 ± 13.4		
G4	178.7 ± 12.7	173.4 ± 22.0	165.3 ± 13.5	177.0 ± 14.4	177.9 ± 11.7*		

* = the mean of body weight differed significantly from the rest of the group means

Parameters	(GO	(G1		G2		G3	(3 4
	Day 0	Day 28	Day 0	Day 28	Day 0	Day 28	Day 0	Day 28	Day 0	Day 28
WBC(M/mm3)	5.9 ± 0.3	5.8 ± 3.7	5.6 ± 1.0	6.2 ± 2.9	5.8 ± 0.5	8.5 ± 3.1	5.3 ± 3.4	10.1 ± 2.4	5.9 ± 0.3	7.7 ± 0.3
Lym (%)	60.8 ± 1.8	57.1 ± 10.5	66.0 ± 18.2	64.7 ± 12.0	59.9 ± 6.	68.8 ± 9.3	61.8 ± 3.1	68.6 ± 14.2	60.8 ± 1.8	68.9 ± 2.3
Mon (%)	12.5 ± 0.5	16.2 ± 4.5	14.4 ± 6.2	17.5 ± 6.1	18.1 ± 1.1	14.6 ± 4.2	18.4 ± 0.1	13.0 ± 3.9	12.5 ± 0.5	14.5 ± 1.2
Neu (%)	6.7 ± 1.3	8.7 ± 4.6	9.6 ± 12.90	7.6 ± 4.3	14.3 ± 2.4	6.5 ± 5.0	6.55 ± 0.2	13.6 ± 5.5	6.7 ± 1.3	10.5 ± 0.9
RBC(M/mm3)	5.8 ± 0.1	6.3 ± 0.0	5.1 ± 0.2	6.1 ± 1.1*	6.5 ± 0.1	5.7 ± 0.6	4.2 ± 1.52	6.12 ± 0.2	5.8 ± 0.1	7.5 ± 0.4
MCV(ft)	54.9 ± 2.5	65.4 ± 1.5	55.7 ± 0.2	55.1 ± 1.2	61.1 ± 0.4	55.5 ± 1.3	45.5 ± 15.5	60.0 ± 2.7	54.9 ± 2.5	62.4 ± 0.8
Hct(%)	31.9 ± 2.0	41.6 ± 1.5	28.7 ± 1.2	33.5 ± 5.3*	39.6 ± 1.0	31.9 ± 4.2	25.1 ± 9.5	36.6 ± 0.4	31.9 ± 2.0	46.0 ± 2.2
MCH(pg)	24.2 ± 0.8	24.0 ± 0.1	25.7 ± 0.3	24.5 ± 1.7	24.2 ± 0.9	25.7 ± 1.5	20.2 ± 7.0	24.8 ± 1.8	24.2 ± 0.8	22.2 ± 0.5
MCHC(g/dL)	44.3 ± 0.5	36.7 ± 0.6	46.5 ± 0.3	34.5 ± 7.7	39.7 ± 1.8	46.5 ± 3.7	33.45 ± 11.3	41.3 ± 1.1	44.3 ± 0.5	35.7 ± 0.3
RDW	11.9 ± 1.3	14.3 ± 0.7	11.4 ± 0.3	11.2 ± 0.5	10.9 ± 0.7	12.1 ± 0.2	10.0 ± 0.6	10.4 ± 0.1	11.9 ± 1.3	10.8 ± 1.0
Hb(g/dL)	14.1 ± 0.7	15.2 ± 0.2	13.2 ± 0.4	14.9 ± 1.4	15.7 ± 0.3	14.7 ± 0.7	11.2 ± 4.2	15.1 ± 0.5	14.1 ± 0.7	16.5 ± 0.6
THR(M/mm3)	1034.5 ± 51.5	2310.5 ± 46.0	1337.5 ± 16.5	1221.0 ± 61.0	1265.5 ± 36.4	1961.0 ± 87.0	1427.0 ± 36.0	997.7 ± 107.2	1034.5 ± 51.5	882.2 ± 12.2
MPV(fl)	9.0 ± 0	8.8 ± 0.0	8.6 ± 0.15	8.7 ± 0.1	8.9 ± 0.2	8.25 ± 0.1	7.6 ± 0.1	6.7 ± 0.2	9.0 ± 0	6.4 ± 0.0
PDW	7.6 ± 0.1	7.7 ± 0.0	8.4 ± 0.25	8.2 ± 0.5	6.85 ± 0.0	8.1 ± 0.0	6.4 ± 0.1	0.7 ± 0.1	7.6 ± 0.1	0.6 ± 0.0

Table 2. Mean hematological parameters before and 28 days after administration chia seed in rats

Table 3. The table indicates the mean plasma glucose levels (Mean \pm S.E) for the control group of rats fed on basal diet treatment group of rats fed on different chia seed feed formulations

Groups	Treatment days						
	0	7	14	21	28		
G0	105.7 ± 4.2	111.5 ± 3.1	118.7 ± 20.5	117.6 ± 2.7	119.9 ± 9.9		
G1	113.5 ± 3.5	122.8 ± 2.6	123.8 ± 7.5	73.5 ± 6.3	61.3 ± 10.9		
G2	116.5 ± 2.0	118.7 ± 4.7	122.1 ± 14.9	102.9 ± 5.6	80.3 ± 12.9		
G3	131.1 ± 1.8	158.1 ± 9.6	131.3 ± 1.1	86.2 ± 7.5	83.1 ± 14.1		
G4	138.5 ± 0.8	110.7 ± 4.7	107.1 ± 4.9	89.9 ± 5.9	80.3 ± 12.9		

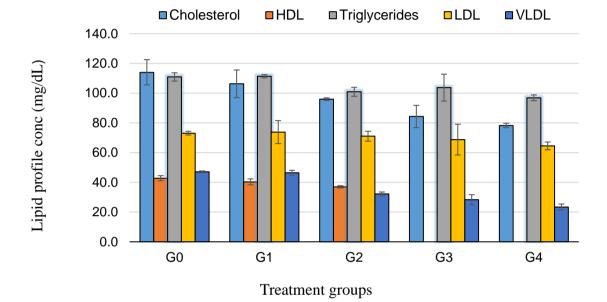


Fig. 1. Plasma lipid profile of rats fed with chia concentrate (Mean ± S.E)

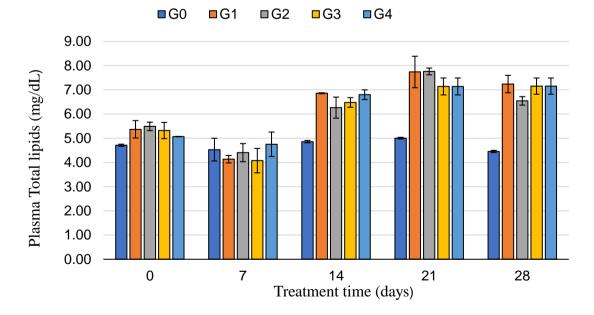


Fig. 2. Plasma total lipids of rats fed with chia concentrate (Mean ± S.E)

Plasma cholesterol concentration: Results in Fig. 1 illustrates that there is significant in all lipid profile in a dose dependent manner (R^2 =99.1; p = 0.000335) compared to the control group GO. The decrease was more in the low density lipoprotein (LDL) and very low density lipiprotein (VLDL).

For plasma total lipids, results indicated there was significant increase (p=0.048) in the level as compared to the control group G0. The level ranged from 4.71 to 7.16 g/dL for G0 to G4 that is, the rats fed with high concentration of chia seed concentrate. For the control groups, the value does not change insignificantly.

4. DISCUSSION

The current study evaluated the effect of chia body seed weight, some selected on hematological and biochemical parameter using rats as model experimental animals. From the study results revealed that the chia seed concentrate had no significant change in body weight of rats fed with chia seed concentrate as compared to the control group. The results concurred with the findings of the study done by Marineli et al. [34], Da Silva et al. [35] and Alamri, [36] who reported insignificant variation of body weight following inclusion of chia seed in feeds. In the current study, the mean body weights for treated rats were lower than the control rats, however the differences were not significant, contrary to observations by other researchers who reported significant weight loss in rats fed on chia seeds. The weight loss in the other studies were linked to the presence of high fibre content in the seeds (more than 80%) [37]. This could also explain the observed weight loss in the rats in this current study. In the current study, dietary supplementation of chia seeds resulted into significant increased in mean values of red blood cell indices such as RBC count, Hb, Hct and. RDW compared to the control group (G0). This could be linked to the high levels of micronutrients such as iron and calcium in the body [38] which are among factors promoting blood formation. The increased level of white blood cell indices observed such as WBC, lymphocytes and monocytes was the indication that chia seeds are capable of boosting the immune system [39,7,40].

All these results are in agreement with the study done by Alarcon et al. [41] Alagawany et al. [42] Mihafu et al. [43]. who reported improvement of hematological markers following dietary supplementation with chia seed in different

animals. The decreased levels of platelets in blood could be due to effect of omega 3 fatty acids which is the important in reducing the content of arachidonic acid in membrane phospholipids of platelets, endothelial cells, and inflammatory cells. This reduces the amount of pro-inflammatory factors that are derived from arachidonic acid. Also Alpha-linoleic acid is found to down-regulate the activity of a nuclear factor NF-KB which plays an important role in generating inflammatory responses which is known to cause cardiovascular diseases [10,44,43]. Alpha-linolenic acid decreases the risk of thrombosis by inhibiting platelet aggregation. They act upon the m RNA expression of the growth factors of platelets, thus stop abnormal growth and proliferation of platelets and play a key role in preventing formation of clots and plaques in the arteries thus elp to prevent cardiovascular diseases [10,44,43]. The plasma glucose and lipid lowering effect of chia seeds have been reported in different animal models as reported from a study by Gian et al. [44]. Diyah, [45] and Radica [46] indicates that the lowereing of blood sugar and lipids by chia seeds is attributable to their high fiber content, and antioxidants present in the chia seed flavonoids. They further indicate that the fibre contents of the seeds cannot be broken down or absorbed. The fibres binds glucose molecules thus preventing it from rapid absorbtion from the gut, a fact that prevents glucose spikes in the blood that can culminate to high blood sugar, diabebetes and elevated high plasma lipids levels.

5. CONCLUSION

Findings in this study illustrated that chia seeds can be the good candidate for controlling body weight, blood glucose, lipid profile and improving hematological parameters in animals. This which are marked as good indicators for controlling metabolic diseases such as as type 2 diabetes mellitus, obesity and cardiovascular diseases are increasing and causes death of both livestock and humans. Further researches are necessary for more confirmation of chia seed on their mode of action and toxicity in animal body.

CONSENT

It is not applicable.

ETHICAL CLEARANCE

The study and all procedures involving the animals were conducted with strict adherence to

guidelines and procedures reviewed and approved by the Institutional Ethical Clearance Committee from the Directorate of the Post graduate, Research and Consultancy Committee of Sokoine University of Agriculture.

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COMPETING INTERESTS

Authors have declared that no competing interests exist.

REFERENCES

- World health organization, noncommunicable diseases progress monitor 2021. Available:https://www.who.int/newsroom/fact-sheets/detail/noncommunicablediseases Accessed:13/11/2023; 2021.
- Dalle GR, Calugi S, Centis E, Marzocchi R, El Ghoch M, Marchesini G. Lifestyle modification in the management of the metabolic syndrome: achievements and challenges. Diabetes Metabolic Syndrome and Obesity. 2010;2(3):373-85.
- 2. Syed IUB. Diet, physical activity, and emotional health: what works, what doesn't, and why we need integrated solutions for total worker health. BMC Public Health. 2020;20:152.
- Scott MG, Barbara H, Sidney CS, James IC, Richard AK. Clinical management of metabolic syndrome: Report of the american heart association/national heart, lung, and blood institute/american diabetes association conference on scientific issues related to management. 2004;09:551–556.
- 4. Melkersson K, Dahl ML. Adverse metabolic effects associated with atypical antipsychotics: literature review and clinical implications. Drugs. 2004;64(7): 701-23.
- Mazereel V, Detraux J, Vancampfort D, van Winkel R De Hert M. Impact of psychotropic medication effects on obesity and the metabolic syndrome in people with serious mental illness. frontiers in endocrinology. 2020;11:573479.
- Graf BL, Raskin I, Cefalu WT, Ribnicky DM. Plant-derived therapeutics for the treatment of metabolic syndrome. current

opinion in investigational drugs. 2010; 11(10):1107-15.

- Khalid W, Arshad MS, Aziz A, Rahim MA, Qaisrani TB, Afzal F, Ali A, Ranjha MMAN, Khalid MZ, Anjum FM. Chia seeds (*Salvia hispanica* I.): a therapeutic weapon in metabolic disorders. food science & nutrition. 2022;11(1):3-16.
- 8. Pascaline O, Jessica B, Sumera A, Nandni K, Arturo A, R'kia M, Lubna S, Savan P, Brionna H, Faaeiza K, Melany P, Parakh B, Sayeed A, Mohd S. 6-Herbal medicines for the treatment of metabolic syndrome, editor(s): Maryam Sarwat, Hifzur Siddique, Herbal Medicines, Academic Press.2022:139-191.
- Ullah R, Nadeem M, Khalique A, Imran M, Mehmood S, Javid A, Hussain J. Nutritional and therapeutic perspectives of chia (*Salvia hispanica* I.): A review. Journal of Food Science and Technology. 2016;53(4):1750-1758.
- Vuksan V, Jenkins AL, Brissette C, Choleva L, Jovanovski E, Gibbs AL, Bazinet RP, Au-Yeung F, Zurbau A, Ho HVT, Duvnjak L, Sievenpiper JL, Josse RG, Hanna A. Salba-chia (*Salvia hispanica* I.) in the treatment of overweight and obese patients with type 2 diabetes: a double-blind randomized controlled trial, nutrition, metabolism and cardiovascular diseases. 2017;27(2):138-146.
- Sara M, Ewa S, Halina E, Agnieszka S. Health-promoting approaches of the use of chia seeds, journal of functional foods. 2023;103:105480
- 12. Masood MB. Chia seeds as potential nutritional and functional ingredients: A Review of Their Applications for Various Food Industries. Journal of Nutrition Food Science and Technology. 2022;4(1):1-14.
- 13. 14.Byars JA, Singh M. Properties of extruded chia–corn meal puffs. lwt-food science and technology. 2015;62(1):506-510
- 14. Avila-Sosa R, Palou E, López-Malo A. Essential oils added to edible films. in essential oils in food preservation, flavor and safety. 2016:149-154. Academic Press.
- 15. Benjamin EJ, Virani SS, Callaway CW, Chamberlain AM, Chang AR, Cheng, de Ferranti SD. Heart disease and stroke statistics-2018 update: a report from the American Heart Association. Circulation. 2018;137(12):e67

- 16. Goyat J, Passi SJ, Suri S Dutta H. Development of chia (*Salvia hispanica*, I.) and quinoa (*Chenopodium quinoa*,I.)seed flour substituted cookies-physicochemical, nutritional and storage studies. current research in nutrition and food science journal. 2018;6(3):757-769
- Ixtaina VY, Nolasco SM, Tomás MC. Physical properties of chia (Salvia hispanica L.) seeds. Industrial Crops and Products. 2008;28:286–293
- Zia-ud D, Mukhtar A, Hidayat Ullah, Dean S, Bin X, Haiteng L, Chaogeng X. Nutritional, phytochemical and therapeutic potential of chia seed (*Salvia hispanica* L.). A mini-review, Food Hydrocolloids for Health. 2021;1:100010
- 19. 20.Vanesa YI, Susana MN, Mabel CT. Physical properties of chia (*Salvia hispanica* L.) seeds, Industrial Crops and Products. 2008;28(3):286-293.
- Kaur N, Chugh V, Gupta AK. Essential fatty acids as functional components of foods- A review. Journal of Food Science and Technology. 2014;51(10):2289-303.
- Dimitra K, Ulrika R, Paul C, Petros K, Leonidas M, Alok P. Benefits of supplementation with microbial omega-3 fatty acids on human health and the current market scenario for fish-free omega-3 fatty acid, Trends in Food Science & Technology. 2023;136:169-180.
- 22. Nadeem M, Imran M, Taj I. Omega-3 fatty acids, phenolic compounds and antioxidant characteristics of chia oil supplemented margarine. Lipids, Health and Diseases. 2017;16:102
- 24.Favati F. Nutritional quality ofseeds and leaf metabolites of Chia (*Salvia hispanica* L.) from Southern Italy. European Food Research and Technology. 2015;241:615-625.
- María del RF, María EO, Victoria A, María E, D'Alessandro. Salvia hispanica L. (chia) seed improves skeletal muscle lipotoxicity and insulin sensitivity in rats fed a sucroserich diet by modulating intramuscular lipid metabolism, Journal of Functional Foods. 2020;66:103775.
- 25. Alamri E. The influence of two types of chia seed on some physiological parameters in diabetic rats. International Journal of Pharmaceutical Research & Allied Sciences. 2019;8(3):131-136
- 26. Toscano LT, da Silva CS, Toscano LT, de Almeida AE, Santos Ada C, and Silva AS. Chia flour supplementation reduces blood

pressure in hypertensive subjects.Plant Foods for Human Nutrition. 2014;69(4): 392–398.

- 27. Mariana G, Thaisa AV, Cintia TS, Andressa de A, Renata LT, Elvira Gonzalez de M, Hercia SDM, Digested protein from chia seed (*Salvia hispanica* L) prevents obesity and associated inflammation of adipose tissue in mice fed a high-fat diet, Pharma Nutrition. 2022;21:100298
- 28. Lucas-Gonzalez Pellearini Μ. R. Fernández-López J, Ricci A, Pérez-Álvarez Sterzo CL. Viuda-Martos Μ. JA, Bioaccessibility of polyphenolic compounds of six quinoa seeds during in vitro gastrointestinal digestion. Journal of Functional Foods. 2017;38:77-88.
- 29. Joaquim C, Carolina P, Andrea A, Ana H, and Ana A. Impact of Processing and Intestinal Conditions on in Vitro Digestion of Chia (*Salvia hispanica*) Seeds and Derivatives, Foods. 2020; 9(3):290
- Parasuraman S, Raveendran R, Kesavan R. Blood sample collection in small laboratory animals. Journal of Pharmacology and Pharmacotherapeutics. 2010;1(2):87-93.
- 31. Trinder P. Determination of Glucose in Blood Using Glucose Oxidase with an Alternative Oxygen Acceptor. Annals of Clinical Biochemistry.1969;6(1):24-27.
- 32. IBM Corp. IBM SPSS Statistics for Windows, Version 28.0. Armonk, NY: IBM Corpz Released; 2021.
- Marineli R, da S, Lenquiste SA, Moraes EA, & Marostica MR. Antioxidant potential of dietary chia seed and oil (*Salvia hispanica* L.) in diet-induced obese rats. Food Research International. 2015;76: 666–674.
- 34. Da Silva, Bárbara P, et al. Chia seed shows good protein quality, hypoglycemic effect and improves the lipid profile and liver and intestinal morphology of wistar rats. Plant Foods for Human Nutrition. 2016;71:225-230.
- 35. Alamri E. The influence of two types of chia seed on some physiological parameters in diabetic rats. International Journal of Pharmaceutical Research and Allied Sciences. 2019; 8(3):131-136.
- 36. Knez HM, Ivanovski M, et al. Chia. Seeds (*Salvia hispanica* L.): An overview-phytochemical profile, isolation methods, and application. Molecules. 2019; 25(1):11.
- 37. Amato M, Caruso MC, Guzzo F, Galgano F, Commisso M, Bochicchio R, Labella

RFD & Favati F. Nutritional quality of seeds and leaf metabolites of Chia (*Salvia hispanica* L.) from Southern Italy. European Food Research and Technology. 2015;241:615-625.

- Monteomo GF, Kamagate A, Yapo AP. Effects of metabolic syndrome on blood cells to wistar rats. Journal of Diabetes, Metabolic Disorders & Control. 2018;5(6):222–225.
- 39. Aparna AR, Abhishek DT, TK, Kanti PS and Sanjay KSP. Nutritional and Functional New Perspectives and Potential Health Benefits of Quinoa and Chia Seeds. Antioxidants, 2023;12(7):1413.
- Alarcon G, Sierra L, Roco J, 40. Van Nieuwenhove C, Medina A, Mirta M, Jere S. Effect of cold pressed chia seed oil intake on hematological and biochemical biomarkers in both normal and hypercholesterolemic rabbits. Plant Foods for Human Nutrition. 2022;78(1): 1-15.
- Mahmoud A; Mohammed N; Al-Abdullatif A; Rashed AA; Mahmoud MA; Fayiz MR. Impact of dietary cold-pressed chia oil on growth, blood chemistry, haematology, immunity and antioxidant status of growing

Japanese quail Italian Journal of Animal Science. 2020;19(1):896-904.

- 42. Mihafu FD, Kiage BN, Kimang'a AN, Okoth JK. Effect of chia seeds (*Salvia hispanica*) onpostprandial glycaemia, body weight and hematological parameters in rats fed a high fat andfructose diet. International Journal of Biological and Chemical Sciences. 2020;14:1752-1762.
- 43. Prathyusha P, Kumari BA, Suneetha WJ, Srujana MNS. Chia seeds for nutritional security. Journal of Pharmacognosy and Phytochemistry. 2019;8(3): 2702-2707.
- 44. Gian CT, Domenico C, Giuseppe B, Maria D, Roberto C, Ettore N. Plasma lipid lowering effect by a novel chia seed based nutraceutical formulation, Journal of Functional Foods. 2018;42:38-46
- 45. Diyah CAK. Kadar Glukosa Darah dan Malondialdehid Ginjal Tikus Diabetes yang Diberi Latihan Fisik. Muhammadiyah Journal of Nursing. 2014;109–116.
- 46. Radica DR, Jakovljevic, Oxidative Stress in Training, Overtraining and Detraining: from Experimental to Applied Research Experimental and Applied Biomedical Research (EABR). 2016;17(4):343-348.

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