

Asian Journal of Research in Biological and Pharmaceutical Sciences

Journal home page: www.ajrbps.com

<https://doi.org/10.36673/AJRBPS.2021.v09.i03.A13>



BIOCHEMICAL STUDY ON THE ROLE OF CURCUMIN IN MALE RABBITS

Sundis A. A. Albera*¹, Fayrouz A. Khaled¹, Marwa J. Said¹

¹*Department Chemistry, Faculty of Science, Omar Al-Mokhtar University, El -Beida-Libya.

ABSTRACT

Turmeric (*Curcuma longa*) may be a therapeutic plant broadly utilized and developed in tropical districts. We have therefore investigate the effects of either 15mg/kg B.W. doses of curcumin on the levels of biochemical parameters in male New Zealand White rabbits. Animals were orally given 15mg/kg B.W. doses of curcumin. The tried measurements were given to rabbits each other day for 6 weeks. Treatment with curcumin caused significant ($P < 0:05$) decrease in the concentrations of glucose, plasma asparatate aminotransferase (AST), alanine aminotransferase (ALT), alkaline phosphatase (ALP) activities, gamma glutamyl transferase activity (γ -GT), Plasma thiobarbituric acid-reactive substances (TBARS) and bilirubin. Contrariwise, the total protein (TP) was significantly increased in plasma. Concentrations of urea. But, creatinine, cholesterol, triglycerides (TG), low-density lipoprotein (LDL) and very low density lipoprotein (VLDL) were significantly decreased, while high density lipoprotein (HDL) was increased. The aim of this study was to evaluate the potential of increasing levels of turmeric powder as phytogetic additives on serum biochemical metabolites and antioxidant enzyme activities in growing rabbits.

KEYWORDS

Curcumin, Biochemistry, Enzyme activities and Rabbits.

Author for Correspondence:

Sundis A. A. Albera,
Department Chemistry,
Faculty of Science, Omar Al-Mokhtar University,
El -Beida-Libya.

Email: fayalzobair@yahoo.com

INTRODUCTON

Turmeric (*Curcuma longa*) may be a restorative plant broadly utilized and developed in tropical locales. Plant extricates were found to have antifungal, imunomodulatory and antioxidative^{1,2} as well as antimutagenic activities³. Turmeric powder is a rich source of beneficial phenolic compounds: the curcuminoids, where three main curcuminoids, curcumin, demethoxycurcumin and bisdemethoxy curcumin⁴ have been isolated from turmeric. Supplementation of turmeric powder at 0.20 and 0.40g/kg to the commercial eat less for rabbits emphatically influenced the body weight pick up in July – September

rabbit does⁵. Turmeric is an Indian rhizomatous herbal plant (*Curcuma longa*) of the ginger family (Zingiberaceae) of well-known medical benefits^{6,7}. Figure No.1 shows *Curcuma longa*. The therapeutic benefits of turmeric might be ascribed to the nearness of active standards called curcuminoids. One of the foremost curcuminoids is curcumin, which could be a little atomic weight polyphenolic compound and lipophilic in nature, subsequently insoluble in water additionally in ether but solvent in ethanol, dimethylsulfoxide, and other natural solvents⁸.

Curcumin is stable at the acidic pH of the stomach⁹. The other constituents present are volatile oils including tumerone, atlantone and zingiberone and sugars, proteins and resins⁷. The active constituent of turmeric- curcumin is isolated from *curcuma longa* and it provides colour to turmeric. Such bioactive component has been thoroughly investigated¹⁰. Curcumin (1, 7-bis (4-hydroxy-3-methoxyphenyl)-1, 6-heptadiene-3, 5-dione) is also called diferuloylmethane¹¹.

On the other hand¹², famous that dietary consideration of turmeric powder at 0, 0.15 and 0.30% had no advantageous impacts on blood parameters and meat characteristics of developing rabbits raised beneath summer stretch. Be that as it may, data around ideal level of garlic and turmeric powder as development promoters and normal cancer prevention agents in developing rabbit is rare. Enhanced concentration of serum AST and ALT are used as indicators of liver damage¹³, the activities of AST and ALT reduced in rabbit fed diet supplemented with turmeric compared to the control group suggesting the hepatoprotective effect of turmeric. Researchers investigate the protective effects of curcumin on experimentally induced hepatotoxicity, and cardio toxicity using various animal models with biochemical parameters like serum marker enzymes and antioxidants in target tissues. The increased relative weight of liver and heart in liver injury and isoproterenol induced cardiac necrosis were also reduced by Curcumin treatment. Elevated serum marker enzymes, aspartate aminotransferase (AST), alanine aminotransferase (ALT) and alkaline phosphatase (ALP) in edematous, granulomatous, liver and heart tissues during liver injury and cardiac necrosis,

respectively. The study demonstrated the *in vitro* and *in-vivo* protective effect of curcumin on experimentally induced hepatotoxicity and cardiotoxicity in rabbits¹⁴.

Within the current explore, bolstering the creatures on turmeric supplemented eat less progressed the hepatic antioxidant enzymes and diminished the MDA concentration in comparison with the control gather. The same results were reported by¹⁵ who found that the intake of herbs resulted in an increase in serum antioxidant enzyme activities and a decrease in MDA concentration.

Our discoveries too upheld by¹⁶ who detailed that turmeric extricates can rummage free radicals, increment antioxidant proteins, and repress lipid peroxidation, though turmeric extricate (1.66mg/kg of body weight) was given to rabbits encouraged a high-fat count calories, oxidation of erythrocyte layers was found to be essentially lower than that in films of control creatures. Grass "metalloprotein chemical" is the primary chemical contributed within the antioxidant defense framework. GSH-Px "seleno chemical" catalyses the reaction of hydro peroxides with diminished glutathione to make glutathione disulphide. Subsequently, hoisted levels of these chemicals may make strides the consistent state of antioxidant framework of rabbits. The concentration of liver MDA is an pointer for assessing antioxidant frameworks. Compared with the control bunch, creatures encouraged 2, 4 or 6g/kg of turmeric had significantly lower liver concentration of MDA common added substances as turmeric can be connected within the future to make strides the wholesome quality of creature meat. It seems that turmeric supplementation to control diet were effective in enhancing the antioxidant ability of animals. Since, turmeric is a rich source of beneficial phenolic compounds, the curcuminoids having strong antioxidant activity^{4,17}. Found that curcumin supplementation restrained lipid peroxidation in rabbit liver microsomes, erythrocyte films and brain homogenates. Besides, lower helplessness of LDL to oxidation¹⁸. Based on these findings, we state that turmeric might play an important role as an exogenous antioxidant and could also be applicable as a protective agent against tissue damage.

MATERIAL AND METHODS

Curcumin was purchased from public market for medicinal herbs in Al-Bayda city. All other chemicals utilized within the explore were of expository review. Mature male New Zealand White rabbits age of 6 months were used. Animals were individually housed in cages and weighed weekly throughout 6-week experimental period. Feed and water were provided ad libitum. Rabbits fed pellets which consisted of 30% berseem (*Trifolium alexandrinum*) hay, 25% yellow corn, 26.2% wheat bran, 14% soybean meal, 3% molasses, 1% CaCl₂, 0.4% NaCl, 0.3% mixture of minerals and vitamins, and 0.1% methionine. The vitamin and mineral premix per kg contained the following IU/gm for vitamins or minerals: vit A-4000,000, vit D3-5000, 000, vit E-16,7g, K0.67g, vit B1-0.67g, vit B2-2g, B6-0.67g, B12-0.004g, B5-16.7g, Pantothenic acid-6.67g, Biotein-0.07g, Folic acid-1.67g, Choline chloride 400g, Zn-23.3g, Mn-10g, Fe-25g, Cu-1.67g, I-0.25g, Se-0.033g, and Mg-133.4g (Rabbit premix produced by Holland Feed Inter. Co.). The chemical analysis of the pellets¹⁹ showed that they contained 15.8 % crude protein, 11.3% crude fiber, 3.7% ether extract, 7.2% ash, 92.9% organic matter and 62.4% nitrogen free extract % as DM basis. Ten develop male rabbits were arbitrarily separated into couple rise to bunches (each five rabbits): Bunch I: Rabbits were utilized as control and gotten an comparable volume of the vehicle (corn oil) alone by oral gavage daily for 6 successive weeks. Group II: Rabbits were treated with curcumin. Curcumin was given daily by gavage at a dose of 15mg/kg B.W¹⁹ which dissolved in corn oil for 6 successive weeks.

At the conclusion of the exploratory period, all rabbits were weighed at that point yielded beneath ether anesthesia. Blood tests were collected in clean dry centrifuge tubes. Plasma was separated by centrifugation at 3000rpm for 10 minutes and then quickly frozen at -20°C for biochemical parameters analysis.

Stored plasma samples were analyzed for total protein (TP) by the Biuret method according to²⁰. Plasma glucose, urea and creatinine concentrations were measured by the method of²¹⁻²³, respectively. Plasma total bilirubin was measured using the

method of²⁴. Plasma concentrations of cholesterol and triglycerides (TG) were determined according to the methods of²⁵⁻²⁷ respectively. High-density lipoprotein (HDL) was determined according to the methods of²⁸. Low-density lipoprotein (LDL) was determined by the calculation (cholesterol-(TG/5+HDL). Very low-density lipoprotein (VLDL) was calculated by dividing the values of TG by factor of 5.

The activities of plasma aspartate transaminase (AST; EC 2.6.1.1) and alanine transaminase (ALT; EC 2.6.1.2) were assayed by the method of²⁹. Alkaline phosphatase (AIP; EC 3.1.3.1) activity was determined in plasma according to the method of³⁰. Plasma thiobarbituric acid-reactive substances (TBARS) were measured by the strategy of Tappel and Zalkin³¹.

RESULTS AND DISCUSSION

Table No.1 showed the overall means of the activities of aspartate transaminase (AST), alanine transaminase (ALT), alkaline phosphatase (AIP), gamma glutamyl transferase activity (γ -GT), Lactate dehydrogenase (LDH), Plasma thiobarbituric acid-reactive substances (TBARS) and bilirubin in blood plasma as affected by treatment with curcumin throughout the 6-week experimental period. Treatment with curcumin resulted in significant ($P < 0.05$) decrease in the activities of blood plasma AST and ALT, (γ -GT), LDH, TBARS and bilirubin while ALP were significantly ($P < 0.05$) increased compared with control group. Table No.2 the mean values of plasma total protein (TP), albumin (A), globulin (G), bilirubin, urea, creatinine and glucose after 6-week experimental period are shown in Table No.2. Treatment with curcumin increase plasma levels of TP, A and G, while decrease glucose, bilirubin, urea and creatinine in male rabbits. Tables 3 illustrated the effect of curcumin on the levels of total cholesterol (TC), triglyceride (TG), very low-density lipoprotein, high and low-density lipoprotein-cholesterol (HDL-c and LDL-c) in blood plasma of male rabbits. The levels of, TC, TG, and LDL-c were significantly ($P < 0.05$) decreased, while HDL-c, were significantly ($P < 0.05$) increased in plasma of rabbits treated with curcumin as compared with control group.

Values are means \pm SEM of 5 rabbits in each group. Mean with different letters (a- d) are significantly difference ($p \leq 0.05$) at same raw. Mean with the same letters (a-d) are non-significantly difference ($p \geq 0.05$).

AST, aspartate amino transferas; ALT, alanin amino transferas; ALP, alkline phosphatase; γ -GT, gamma glutamyl transe activity; Lactate dehydrogenase, TBARS, thiobarbituric acid-reactive substances, Cho., cholosterol; TG, triglycerides; HDL, high density lipoprotein; LDL, low density lipoprotein.

Discussion

The protective effect of curcumin may be explained by the fact that it prevents cellular damage occurring as a result of oxidative stress³². Administration of turmeric resulted in slight decrease level of liver enzymes as compared to sodium nitrite treated group indicate to ameliorative effect of turmeric ,These results are in harmony with^{33,34}, they stated that curcumin administration prevented ALT and AST increases and improved liver function.

The obtained results were in agreement with³⁵ who found that triglycerides, total cholesterol and low density lipoprotein concentrations were linearly and quadratically decreased with increasing the dietary proportion of turmeric. Also³⁶, clearly found that triglycerides and total cholesterol were significantly decreased with increasing turmeric levels to 7g/kg diet. The reduction of lipids profile (triglycerides, total cholesterol and low density lipoprotein concentrations) may be due to curcumin that enhances bile production and hence lipid digestion^{37,38}. Indicated that liver triacylglycerol and cholesterol concentrations were significantly lower in rats fed curcumin than in control animals. Plasma triacylglycerol in the very low-density lipoproteins fraction were also lower in curcumin fed rats. Hepatic acyl-CoA oxidase activity of the curcumin group was significantly higher than that of the control. The obtained results in the present study contrast with the results reported by³⁹ who demonstrated that turmeric supplementation into the basal diets of broilers significantly increased total cholesterol and HDL cholesterol and decreased LDL-cholesterol, but did not affect total-triglycerides.

Table No.1 presents the results on the effects of different supplementations on serum malondialdehyde concentrations of rabbits. Turmeric supplemented diet reduced the MDA concentration in comparison with the control group⁴⁰. Reported that supplementation with Curcuma longa reduces oxidative stress and attenuates the development of fatty streaks in rabbits fed a high cholesterol diet. Such antioxidant effects would be expected to improve the health of rabbits. From these results, it can be stated that supplementation with phytogetic additives as turmeric could be applied in the future to improve the nutritional quality of rabbit meat. It seems that turmeric supplementation to basal control diet was effective in enhancing the antioxidant ability of rabbits. These effects are due to the antioxidant property of turmeric^{4,18}. Found that curcumin supplementation inhibited lipid peroxidation in rat liver microsomes, erythrocyte membranes and brain homogenates. Moreover, it lowered susceptibility of LDL⁴¹. Found that curcumin protects against chemical toxicity.

Table No.1: Changes in the biochemical parameters and the level of thiobarbituric acid-reactive substances (TBARS) during treatment of male rabbits with 100mg/kg doses of curcumin

S.No	Parameters	Animal Groups	
		Control	Curcumin
1	AST (U/L)	43.30±1.032 ^a	39.19±3.443 ^a
2	ALT (U/L)	45.35±0.706 ^a	39.18±2.809 ^a
3	ALP (U/L)	144.89±3.769 ^a	153.48±6.941 ^a
4	γ-GT (U/L)	7.13±0.029 ^a	6.60±0.081 ^b
5	Bilirubin (mg/dl)	1.56 ±0.009 ^a	1.47 ±0.024 ^b
6	TBARS (nmol/ml)	0.701 ± 0.019 ^a	0.602 ± 0.039 ^a
7	TP (g/dl)	6.83±0.021 ^a	6.98 ± 0.119 ^a
8	Glucose (mg/dl)	115.36 ± 0.510 ^a	113.43 ± 1.504 ^a
9	Urea (mg/dl)	37.97±0.277 ^a	34.05± 0.710 ^b
10	Creatinine (mg/dl)	0.63 ±0.005 ^a	0.53±0.030 ^b
11	Cho	120.40 ± 0.494 ^a	118.05 ±0.706 ^b
12	TG	55.75 ± 0.693 ^a	54.22 ±1.078 ^b
13	HDL	57.47± 0.618 ^b	61.31 ± 7.25 ^a
14	LDL	59.33 ± 0.840 ^a	54.93 ± 0.693 ^b

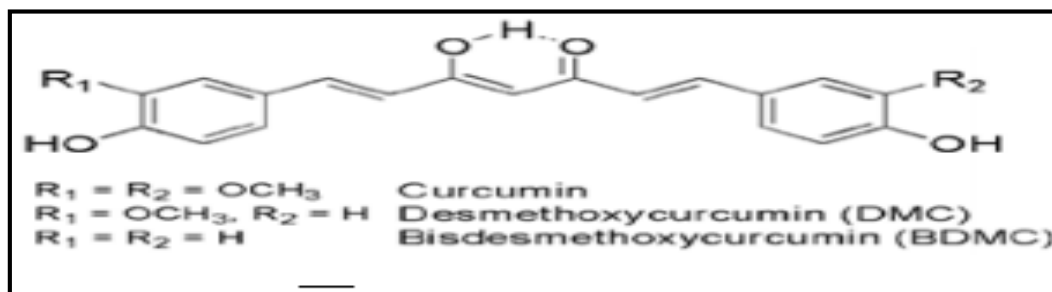


Figure No.1: Chemical structures of curcumin

CONCLUSION

Based on experiment findings, we state that turmeric could have beneficial effects on biochemical parameters and may be play an important role as an exogenous antioxidant.

ACKNOWLEDGEMENT

The authors wish to express their sincere gratitude to Department Chemistry, Faculty of Science, Omar Al-Mokhtar University, El -Beida-Libya for providing necessary facilities to carry out this research work.

CONFLICT OF INTEREST

We declare that we have no conflict of interest.

BIBLIOGRAPHY

1. Farag M R, Alagawany M M, Dhama K. Antidotal effect of turmeric (*Curcuma longa*) against endosulfan-induced cytogenotoxicity and immunotoxicity in broiler chicks, *International Journal of Pharmacology*, 10(8), 2014, 429-439.
2. Alagawany M, Ashour E A, Reda F M. Effect of dietary supplementation of garlic (*Allium sativum*) and turmeric (*Curcuma longa*) on growth performance, carcass traits, blood profile and oxidative status in growing rabbits, *Annals of Animal Science*, 16(2), 2016, 489-505.
3. Soni K B, Lahiri M, Chackradeo P, Bhide S V, Kuttan R. Protective effect of food additives on aflatoxin-induced mutagenicity and hepatocarcinogenicity, *Can Let*, 115(2), 1997, 129-133.

4. Balasubramanyam M, Koteswari A A, Kumar R S, Mohan V. Curcumin-induced inhibition of cellular reactive oxygen species generation: novel therapeutic implications, *Jour of Bio*, 28(6), 2003, 715-721.
5. Foldesiova M, Balazi A, Chrenek P. The effect of Curcuma longa dried powder in the diet on weight gain of rabbit does, *Slovak Jour of Ani Sci*, 48(1), 2015, 43-48.
6. Panpatil V V, Tattari S, Kota N, Nimgulkar C, Polasa K. *In vitro* evaluation on antioxidant and antimicrobial activity of spice extracts of ginger, turmeric and garlic, *Jourof Phar Phy*, 2(3), 2013, 143-148.
7. Pawar H, Karde M, Mundle N, Jadhav P, Mehra K. Phytochemical evaluation and curcumin content determination of turmeric rhizomes collected from Bhandara District of Maharashtra (India), *Med. Chem*, 4(8), 2014, 588-591.
8. Aggarwal B B, Kumar A, Bharti A C. Anticancer potential of curcumin: Preclinical and clinical studies, *Anti Res*, 23(1/A), 2003, 363-398.
9. Kharat M, Du Z, McClements D J. Physical and chemical stability of curcumin in aqueous solutions and emulsions: Impact of pH, temperature, and molecular environment, *Jou of Agr and Food Ch*, 65(8), 2017, 1525-1532.
10. Hewlings S J, Kalman D S. Curcumin: A review of its effects on human health, *Foods*, 6(10), 2017, 92.
11. Alsamydai A, Jaber N. Pharmacological aspects of curcumin, *Int J Pharm*, 5(6), 2018, 313-326.
12. Basavaraj M, Nagabhushana V, Prakash N, Appannavar M M, Waggmare P, Mallikarjunappa S. Effect of dietary supplementation of Curcuma longa on the biochemical profile and meat characteristics of broiler rabbits under summer stress, *Veterinary World*, 4(1), 2011, 15-18.
13. Ozaki M, Fuchinoue S, Ota K. The *in vivo* cytoprotection of ascorbic acid against ischemia/reoxygenation injury of rat liver, *Ar of Bio and Bioph*, 318(2), 1995, 439-445.
14. Scannell J W, Blanckley A, Boldon H, Warrington B. Diagnosing the decline in pharmaceutical R and D efficiency, *Nature Reviews Drug Discovery*, 11(3), 2012, 191-200.
15. Lin C C, Wu S J, Chang C H, Ng L T. Antioxidant activity of Cinnamomum cassia, *Phytotherapy Research*, 17(7), 2003, 726-730.
16. Wachtel-Galor S, Yuen J, Buswell J A, Benzie I F. Ganoderma lucidum (Lingzhi or Reishi), Herbal medicine: Biomolecular and clinical aspects, *CRC Press/Taylor and Francis*, 2nd Edition, 2011.
17. Reddy A C P, Lokesh B R. Studies on the inhibitory effects of curcumin and eugenol on the formation of reactive oxygen species and the oxidation of ferrous iron, *Molecular and Cellular Biochemistry*, 137(1), 1994, 1-8.
18. Mesa M D, Ramírez-Tortosa M C, Aguilera C M, Gil A. Nutritional and pharmacological effects of Curcuma longa L. extracts, *Recent Research Developments in Nutrition*, 3, 2000, 157-171.
19. Ahmad R M, AL-Hubaity A Y, Alazow N S. The role of Vitamin C on the structural changes of male albino rats kidney induced by tramadol, *Annals of the College of Medicine, Mosul*, 41(1), 2019, 57-62.
20. Armstrong W, Carr C. Physiological chemistry laboratory directions, (*Minneapolis, MN, Burges*), 3rd Edition, 1964.
21. Trinder P. Cited from Chmory Enzymatic glucose reagent set (colorimetric), *Ann, Clin. Biochem*, 6(2), 1969.
22. Patton C J, Crouch S R. Spectrophotometric and kinetics investigation of the Berthelot reaction for the determination of ammonia, *Analytical Chemistry*, 49(3), 1977, 464-469.
23. Henry R, Cannon, D, Winkelman W. Clinical chemistry principals and techniques, *Happer and Row Publishers, New York*, 11th Edition, 1974, 1629.
24. Pearlman F C, Lee R T. Detection and measurement of total bilirubin in serum, with use of surfactants as solubilizing agents, *Clinical Chemistry*, 20(4), 1974, 447-453.
25. Knight J A, Anderson S, Rawle J M. Chemical basis of the sulfo-phospho-vanillin

- reaction for estimating total serum lipids, *Clinical Chemistry*, 18(3), 1972, 199-202.
26. Watson D. A simple method for the determination of serum cholesterol, *Clinica Chimica Acta*, 5(5), 1960, 637-643.
 27. Fossati P, Prencipe L. Serum triglycerides determined colorimetrically with an enzyme that produces hydrogen peroxide, *Clinical Chemistry*, 28(10), 1982, 2077-2080.
 28. Warnick G, Benderson J, Albers J J. Dextran sulfate-Mg²⁺ precipitation procedure for quantitation of high-density-lipoprotein cholesterol, *Clinical Chemistry*, 28(6), 1982, 1379-1388.
 29. Reitman S, Frankel S. A colorimetric method for the determination of serum glutamic oxalacetic and glutamic pyruvic transaminases, *American Journal of Clinical Pathology*, 28(1), 1957, 56-63.
 30. Principato G B, Aisa M C, Talesa V, Rosi G, Giovannini E. Characterization of the soluble alkaline phosphatase from hepatopancreas of *Squilla mantis* L, *Comparative Biochemistry and Physiology Part B: Comparative Biochemistry*, 80(4), 1985, 801-804.
 31. Tappel A L, Zalkin H. Inhibition of lipide peroxidation in mitochondria by Vitamin E, *Ar of Bio and Bioph*, 80(2), 1959, 333-336.
 32. Aly H A, Mansour A M, Abd-Ellah H F, Abdel-Naim A B. Potential testicular toxicity of sodium nitrate in adult rats, *Food and Che Toxi*, 48(2), 2010, 572-578.
 33. Hemeida R A, Mohafez O M. Curcumin attenuates methotrexate-induced hepatic oxidative damage in rats, *Journal of the Egyptian National Cancer Institute*, 20(2), 2008, 141-148.
 34. Hassan S M., Zagloul N F, El-shamy S A. Comparative studies on turmeric and Vitamin C on sodium nitrite treated rats, *Alexandria Jour for Vet Sci*, 56(1), 2018, 56-68.
 35. Alagawany M M, Farag M R, Kuldeep D. Nutritional and biological effects of turmeric (*Curcuma longa*) supplementation on performance, serum biochemical parameters and oxidative status of broiler chicks exposed to endosulfan in the diets, *Asian Journal of Animal and Veterinary Advances*, 10(2), 2015, 86-96.
 36. Hussein S N. Effect of turmeric (*Curcuma longa*) powder on growth performance, carcass traits, meat quality, and serum biochemical parameters in broilers, *Journal of Advanced Biomedical and Pathobiology Research*, 3(2), 2013, 25-32.
 37. Al-Sultan S I, Gameel A A. Histopathological changes in the livers of broiler chicken supplemented with turmeric (*Curcuma longa*), *International Journal of Poultry Science*, 3(5), 2004, 333-336.
 38. Asai A, Miyazawa T. Dietary curcuminoids prevent high-fat diet-induced lipid accumulation in rat liver and epididymal adipose tissue, *The Journal of Nutrition*, 131(11), 2001, 2932-2935.
 39. Emadi M, Kermanshahi H, Maroufyan E. Effect of varying levels of turmeric rhizome powder on some blood parameters of broiler chickens fed corn-soybean meal based diets, *Int. J. Poult. Sci*, 6(5), 2007, 345-348.
 40. Quiles J L, OcHoA J J, Huertas J R. Olive oil and mitochondrial oxidative stress: Studies on adriamycin toxicity, physical exercise and ageing, *Olive Oil and Health*, Oxford: CABI Publishing, 76(4), 2006, 119-151.
 41. Iqbal M, Sharma S D, Okazaki Y, Fujisawa M, Okada S. Dietary supplementation of curcumin enhances antioxidant and phase II metabolizing enzymes in ddY male mice: possible role in protection against chemical carcinogenesis and toxicity, *Pharmacology and Toxicology*, 92(1), 2003, 33-38.

Please cite this article in press as: Sundis A. A. Albera et al. Biochemical study on the role of curcumin in male rabbits, *Asian Journal of Research in Biological and Pharmaceutical Sciences*, 9(3), 2021, 89-95.