

Vitamin C Supplementation Lowered Atherogenic Lipid Parameters Among Oil and Gas Workers Occupationally Exposed to Petroleum Fumes in Port Harcourt, Rivers State, Nigeria

ABSTRACT

Aim: To evaluate the effect of vitamin C supplementation on lipid profile parameters among oil and gas workers occupationally exposed to petroleum fumes through inhalation over a period of 12 months.

Study Design: A total of 150 subjects between the age ranges of 18 to 45 years old were recruited for the study of which 50 subjects were exposed to petroleum fumes and were given vitamin C supplementation (group 1), 50 were also exposed to petroleum without vitamin C (Group 2) and 50 subjects are non-oil and gas workers (control subjects).

Methodology: The Group 1 subjects were orally administered vitamin C supplements of 100mg/day for 60 days before sample collection while Group 2 subjects exposed to petroleum fumes were not given vitamin C or took vitamin during the period of the study. The control subjects were non-oil and gas workers. At the end of the experiment (60 days), blood specimens were collected from group 1, group 2, and the control subjects. Lipid parameters such as cholesterol, triglycerides, high-density lipoprotein, low-density lipoprotein, and very low-density lipoprotein were analysed using enzymatic methods. Statistical analysis was done using GraphPad Prism and results were expressed as Mean±SD.

Results: The results indicated significantly higher values in total cholesterol, triglyceride, low-density lipoprotein, and very low-density lipoprotein in the petroleum fumes exposed subjects compared to control subjects except for high-density lipoprotein which indicated significantly lower values. When those exposed were given vitamin C supplementation, there was a significant reduction in the lipid parameters except HDL-C which indicated a significantly higher value compared with those exposed without vitamin C supplementation administered orally at 100mg for 60 days at $P=0.05$.

Conclusion: The study has shown that occupational exposure to petroleum fumes is associated with dyslipidaemia. However, the use of 100mg of vitamin C daily for 60 days ameliorated the degree of dyslipidaemia associated with occupational exposure to petroleum fumes. Vitamin C could be considered as a preventive means to mitigate or ameliorate cardiovascular risks due to occupational exposure to petroleum fumes and products among workers.

Keywords: Gasoline pump workers, gasoline, lipid profile, vitamin C, anti-oxidant, petroleum fumes, Dyslipidaemia, cardiovascular risk

1. INTRODUCTION

Lipids function as hormones, energy sources, and structural components in cell membranes and are transported in the plasma binding to specific carrier proteins called lipoproteins [1]. Lipid particles such as total cholesterol, triglycerides, high-density lipoprotein, low-density lipoprotein, and very low-density lipoprotein are vital biochemical parameters usually employed as a frontline approach in the screening for abnormalities in lipids metabolism viz-a-viz in disease diagnosis such as cardiovascular diseases [2, 3]. Lipoproteins include chylomicrons, very low-density lipoprotein (VLDL), low-density lipoprotein-cholesterol (LDL-C), and high-density lipoprotein-cholesterol (HDL-C) [2]. Abnormalities in the metabolism of lipids and lipoproteins are linked with the risks of developing atherosclerosis, one of the underlying causes of cardiovascular disorders such as myocardial infarction, cerebrovascular diseases, and peripheral vascular disease [2, 3]. Several factors such as exposure to drugs, chemicals, and lifestyle have been reported to affect lipid levels. Elekima et al. [1], documented that, oral administration of carmoisine (a food

dye) affected the lipid metabolism with a resultant increase in cholesterol level. Exposure to petroleum or petroleum fumes could also serve as a risk for dyslipidaemia.

Petrol is a clear petroleum-derived flammable liquid that is used primarily as fuel in most spark-ignited internal combustion engines[4]. It consists mostly of organic compounds obtained by the fractional distillation of petroleum, enhanced with a variety of additives [5].The hazardous effect of petrol (fuel) on human health has greatly been of public health concern and filling station fuel pump workers are an important group who are at occupational risk to BTX (benzene, toluene, and xylene) compounds which are the main constituents of petrol of which Benzene stands out for its hazardous effects on human health[4, 6, 7]. Keenan et al. [8], documented that that simultaneous exposure to benzene and other aromatic hydrocarbons, such as toluene and xylene, contributes to maximizing benzene toxicity and oxidative stress.

There are health concerns for workers occupationally exposed to benzene [4, 5]. These concerns are linked to the fact that benzene is a well-recognized genotoxic human carcinogen, classified as a Group I chemical by the International Agency for Research on Cancer, and without any known threshold dose [7].Occupational Safety and Health Administration [5] has set a permissible exposure limit of 1 ppm of benzene in air at the workplace during an 8-hour workday, and 40-hour workweek. OSHA, [5], further stated that the short-term exposure limit for airborne benzene is 5 ppm for 15 minutes. The U.S. National Institute for Occupational Safety and Health (NIOSH) in 2004 revised the immediately dangerous to Life and Health (IDLH) concentration for benzene to 500ppm.

Badham et al.[9], documented that the activation of benzene and its metabolites in petroleum products culminates in damage to lipids, proteins, DNA, and carbohydrates through various chemical reactions involving oxidative stress, leading to functional alterations in different tissues. Moro *et al.*,[4] also further reported that are exposed to benzene in petroleum products can lead to non-cancer health effects, such as genotoxicity, hematotoxicity, hepatotoxicity, and nephrotoxicity. The mechanism of action of their toxicity has been linked with the rapid generation of free radicals overwhelming the anti-oxidant capacity of the system leading to peroxidation of lipids and consequently inducing local injury to cell membranes[10, 11].

Free radicals are reactive chemicals that possess the capacity to disrupt cellular activities. These radicals are usually generated during oxidative processes when an atomic or molecular moiety either gains or loses an electron. They could be introduced into the system internally through metabolic processes or externally [12]. Free radicals in required physiological levels in the blood and cellular components are involved in cellular signals and other vital functions [2, 12]. However, when their production overwhelms the anti-oxidant capacity, it becomes pathologic resulting in different disorders affecting the macromolecules and micro-molecules, including DNA, proteins, lipids, proteins, and cell membranes. Free radicals with oxygen as the active agent are referred to as reactive oxygen species (ROS) while those with nitrogen are referred to as reactive nitrogen species (RNS). ROS are the most common type of free radicals produced in living tissue[12]]. The effects of these free radicals are usually counteracted by an antioxidant defense mechanism.

Antioxidants are chemicals whose main function is in the eradication of free radicals, allowing physiologically permissible levels [12]. Antioxidants are produced within the system (endogenous antioxidants) or derived (Exogenous anti-oxidants). Irrespective of the endogenous anti-oxidant generation, the biological system also depends on exogenous sources, mainly, through the diet (fruits, vegetables, and grains) and dietary supplements[12]. Examples of dietary antioxidants include beta-carotene, lycopene, and vitamins A, C, and E. The human body's complex antioxidant defense system

constitutes dietary intakes of antioxidants in the form of fruits and vegetables and endogenous production of antioxidant compounds e.g. glutathione [12].

Vitamin C (Ascorbic Acid) is considered a vitamin with a significant anti-oxidative effect and by extension could be exceptionally gainful in the therapy of oxidative-induced organ damage. Studies by George-Opuda et al., [13], revealed that Vitamin C reduced the impact of gasoline exposure in rats. Vitamin C is a powerful reducing agent that is a water soluble vitamin required for the maintenance of skin and involved in the removal of reactive species. The essence of the study is to assess the influence of Vitamin C on the lipid parameters after exposure to petroleum fumes.

2. Materials and Methods

2.1 Materials

The materials used in this study included vitamin C, a vacutainer, lithium heparin bottles, an automatic pipette, test tubes, and a spectrophotometer.

2.2 Study Design

A total of 150 healthy adults between the ages of 18 to 45 years were recruited for the study. One hundred of the participants are working as fuel attendants at gas stations exposed to petroleum fumes. Of these 100 exposed subjects, 50 subjects (Group 1) were given 100mg of vitamin C supplements daily for 60 days while the other 50 (Group 2) were not given vitamin C or took vitamins during the study period. The control subjects were non-oil and gas workers. After 60 days, blood samples were collected from the control group, and exposed groups for laboratory analyses after 12 hours fast.

2.3 Subjects Selection Criteria

2.3.1 Inclusion Criteria

Subjects included in the study were between the ages of 18-45 years. They are non-alcoholics, not on any medication, or chronic diseases. Also, the filling station workers had worked in the fuel station for a minimum of one year.

2.3.2 Exclusion Criteria

Those excluded included alcoholics, those with a history of cardiac or cardiac-related disease, and those on medication. Staff members recently employed or have worked less than 12 months were also excluded.

2.4 Sample Collection and Preparation

Venous blood specimens in fasting states were collected into lithium heparin bottles and centrifuged to obtain plasma. The plasma obtained was used to determine the concentration of cholesterol, triglycerides, and high-density lipoprotein.

2.5 Sample Analysis

Cholesterol and triglycerides were determined using the enzymatic methods. High-density lipoprotein (HDL) cholesterol was analysed using the precipitation method while LDL cholesterol levels were calculated using Friedwald's equation as described by Friedwald et al. [14].

Statistical Analysis

Statistical analysis was performed using SPSS. Results were presented as Mean±SD. Student's t-test was used to compare between control and test subjects. Also, One-Way ANOVA (Post Hoc: Tukey's multiple comparative tests) was used to determine the significant differences between Group 1, Group 2, and the control.

3 RESULTS and DISCUSSION

3.1 Result of Lipid Parameters in Subjects Exposed to Petroleum Fumes

The results indicated significantly higher values in total cholesterol, triglyceride, and very low-density lipoprotein in the fuel fumes exposed subjects at $P=.05$ (Table 1). However, high-density lipoprotein indicated higher values in unexposed compared to the exposed subjects at $P=.05$ (Table 1).

Table 1. Results of Lipid Parameters in Subjects Exposed to Petroleum Fumes

Parameters	Exposed (Test)	Unexposed (Control)	P value	Remark
TCHOL(mmol/L)	4.629±0.449	3.980±0.555	<0.0001	S
TRIG (mmol/L)	1.196±0.301	0.908±0.505	0.0008	S
HDL (mmol/L)	0.936±0.181	1.034±0.211	0.0155	S
LDL (mmol/L)	3.002±0.509	2.635±0.500	0.0004	S
VLDL (mmol/L)	0.520±0.154	0.408±0.232	0.0057	S

Keys: HDL= High-density lipoprotein, TCHOL = Total cholesterol, LDL= Low-density lipoprotein, VLDL= Very low-density lipoprotein. S=Significant at $P=.05$

3.2 Result of Lipid Parameters in Control Subjects, and Subjects Exposed Petroleum fumes with and without vitamin C supplementation

The results indicated significantly higher values in subjects exposed to petroleum fumes without Vitamin C supplementation compared to the subjects exposed to petroleum fumes given 100mg of Vitamin C supplementation for 60 days. It was further observed that there were significant differences in the values of the lipid parameters considered when workers exposed to petroleum fumes were compared with control subjects except in HDL-C where those that were exposed and had vitamin C supplementation for 60 days had higher values compared to control and exposed groups without supplementation at $P=.05$ (Table 2). More so, the results indicated showed that those exposed without vitamin C supplementation had the lowest values of HDL-C compared to control (without vitamin C) and those exposed with vitamin C supplementation at $P=.05$.

Table 2: Result of Lipid Parameters in Control Subjects, Petroleum Fume Exposed-vitamin C Supplementation, and Petroleum Fume Exposed without Vitamin C Supplementation

Parameters	Control	Exposed + Vitamin C (Group 1)	Exposed Without Vitamin C (Group 2)	P value	Remark
T.CHOL(mmol/L)	4.103 ± 0.222 ^a	4.201±0.123 ^a	4.687 ± 0.841 ^b	<0.0001	S
TRIG (mmol/L)	1.004 ± 0.201 ^a	1.210±0.658 ^a	1.496 ± 0.801 ^b	0.0006	S
HDL (mmol/L)	0.936 ±0.181 ^a	1.187 ± 0.194 ^b	0.895±0.222 ^c	0.0166	S
LDL (mmol/L)	2.635 ±0.100 ^a	2.234±0.231 ^a	3.982 ± 0.604 ^b	0.0002	S
VLDL (mmol/L)	0.408 ±0.232 ^a	0.507±0.089 ^a	0.610 ± 0.254 ^b	0.0059	S

Keys: HDL= High-density lipoprotein, TCHOL = Total cholesterol, LDL= Low-density lipoprotein, VLDL= Very Low-density lipoprotein. S=Significant at $P=.05$. PostHoc: Values within the same row with different superscripts differ significantly at $P=.05$

The present study investigated the effect of vitamin C supplementation on the lipid parameters of subjects exposed to petroleum products or fumes for over a year. The significantly higher values observed in the lipid parameters, that is, total cholesterol, triglycerides, low-density lipoproteins, and very-low-density lipoproteins in the fuel pump workers compared to non-fuel pump workers similar to the reports of Uboh et al., [15], who reported an increase in all lipid parameters of rats exposed to petrol & kerosene fumes. However, high-density lipoproteins indicated significantly highest value in those exposed and had vitamin C supplementation compared to control and those exposed without supplementation. This finding is in line with the reports of Festus et al. [16], who reported a significant decrease in HDL-C plasma levels of mechanics occupationally exposed to petroleum products and fumes alongside other lipid particles. In addition, Ubani et al. [17], documented a significant reduction in HDL-C values in albino rats exposed to 0.10%, 0.50%, and 1.00% of gasoline for 14 days. In the same view, Chukwurah et al [18], also observed significantly higher values of HDL-C in rats exposed to premium motor spirit (PMS) for 28 days at 1, 2, and 5 hours daily. However, our finding is contrary to the reports of Mohammed et al. [19], who reported a significant increase in plasma HDL-C levels of individuals exposed to gasoline from different derivatives and who were also smoking. They further stated that smoking exacerbated the increases observed in HDL-C plasma concentration.

Furthermore, the significantly higher values of total cholesterol, triglycerides, low-density lipoproteins, and very low-density lipoproteins in our findings are in line with the documentation of Ugbala and colleagues [20]. They reported a dose-dependent increase in total cholesterol and low-density lipoprotein cholesterol (LDL-C). However, their study reported a significantly lower value of triglyceride concentration which is contrary to our findings. This disparity in our results and that of Ugbala and colleagues [20] could be due to the fasting state in our study (18 hours) prior to blood collection. Ogbavire et al. [21] in their study reported significantly higher plasma levels of cholesterol, triglycerides, and low-density lipoprotein cholesterol in rats when exposed to premium motor spirit (PMS) fumes but documented no significant difference in high-density lipoprotein cholesterol (HDL-C) values. In a similar study, Kapil et al. [22] and Egbuonu et al. [23] observed dyslipidaemia among workers at petroleum depots in Calabar, Nigeria, and South Haryana, India respectively occupationally exposed to PMS products and fumes. The significantly higher values of these lipid parameters indicated that exposure to gasoline vapours could be associated with hyperlipidaemia, and therefore a risk factor for atherogenicity. The increase in LDL-cholesterol and the corresponding decrease in HDL-cholesterol could be a result of gasoline-inducing cellular injury and functional abnormalities in hepatocytes by the process of lipid peroxidation since the liver plays a central role in the maintenance of lipid homeostasis.

However, when Vitamin C supplementation was introduced in an already exposed subject at a dose of 100mg daily for over a period of 60 days, the levels of cholesterol, triglycerides, LDL-C, and VLDL-C were significantly lowered similar to those of the control (non-attendants) and therefore no significant difference between these groups except HDL-C that indicated highest values in those taking vitamin C compared to control and those exposed without vitamin C supplementation. However, those exposed without the use of vitamin C still had significantly higher values of these lipid parameters. The results observed suggest that vitamin C supplementation enhanced the anti-oxidative capacity of the system, therefore involved in the counteracting of free radicals generated via the inhalation of chemicals from petroleum fumes. This finding concurs with the observation of Gaur & Dixit, [24], who reported that the administration of C caused a significant reduction in serum total cholesterol and LDL cholesterol. More so, George-Opuda et al. [13], documented in their work that vitamin C was demonstrated as an effective anti-oxidant, particularly in oxidation-induced circumstances.

Vitamin C has been documented by Pehlivan [25] as a natural antioxidant present in both animals and plants. This vitamin plays a vital role in the body's reduction-oxidation reactions therefore involved in neutralizing the activities of reactive oxygen or nitrogen species produced during metabolism. Pehlivan [25] further documented that vitamin C functions as a cofactor for some enzymes associated with hormone biosynthesis and regenerating other antioxidants, therefore playing a synergic role in anti-oxidation.

Our results by implication suggest that vitamin C at a daily dose of 100mg plays a potential role in mitigating atherosclerotic and cardiovascular risk associated with fuel toxicity through inhalation, **therefore playing a cardio-protective role amongst those exposed to petroleum fumes.** This finding is particularly relevant for individuals exposed to occupational hazards, such as fuel pump workers, who may be at an increased risk of cardiovascular diseases due to occupational exposures. In other words, this study has demonstrated that vitamin C has a protective effect at 100mg daily against induced dyslipidaemia through the inhalation of petroleum fumes among workers exposed occupationally. More so, the result provides additional support for considering vitamin C as a preventive or ameliorative drug for cardiovascular risks among workers exposed to petroleum fumes chronically since vitamin C is generally safe and its use is not associated with side effects when administered within recommended doses. **Sebastian et al. [26], also documented that epidemiological studies indicated that vitamin C-rich diets such as fruits and vegetables are linked with reduced risk of cardiovascular disease, cancer, and other chronic diseases due to their anti-oxidant and pro-oxidant roles.** This safety profile of vitamin C, **cost-effectiveness, availability, and means of administration** makes it a favorable candidate for preventive or ameliorative interventions, especially in occupational settings where exposures to petroleum fumes are unavoidable.

5. CONCLUSION

The study has shown that occupational exposure to petroleum fumes is associated with dyslipidaemia. However, the use of 100mg of vitamin C daily for **60 days** ameliorated the degree of dyslipidaemia associated with occupational exposure to petroleum fumes. Vitamin C could be considered as a preventive means to mitigate or ameliorate the adverse effects of occupational exposure to petroleum fumes and products among workers.

Ethical Approval:

As per international standards or university standards written ethical approval has been collected and preserved by the author(s).

Consent

As per international standards or university standards, respondents' written consent has been collected and preserved by the author(s).

Funding

This research work was financially supported by the authors.

Competing Interests

The authors have declared that no competing interests exist.

References

1. Elekima I, Nwachuku, EO, Ben-Chioma AE. Effect of tartrazine orally administered on the lipid profile of albino rats. *European Journal of Pharmaceutical and Medical Research*. 2017;4(7): 164-167
2. Elekima I. Effect of carmoisine orally administered on lipid parameters of albino rats. *International Journal of Science and Research (IJSR)*. 2016;5(9): 861 -864

3. Elekima I, Onwuli D, Obisike UA, Aleru CP, Christian SG, Osiagor J. Effect of hibiscus sabdariffa extract orally administered on lipid parameters of albino rats. *Journal of Medical Science and Clinical Research*. 2016; 4 (2): 9529-9533]
4. Moro AM, Brucker N, Chara, MF, Baierle M, Sauer E, Goethel G. Biomonitoring of gasoline station attendants exposed to benzene: effect of gender. *Mutation Research*, 2017; 813: 1-9.
5. OSHA. Chemical Sampling Information Benzene. 2011) Retrieved from Osha.gov. Retrieved on 2011-11-23.
6. Dougherty D, Garte S, Barchowsky A, Zmuda J, Taioli E. (2008). NQO1, MPO, CYP2E1, GSTT1 and GSTM1 polymorphisms and biological effects of benzene exposure —a literature review. *Toxicology Letters*. 2008; 182(13), 7-17.
7. Carrieri M, Spataro G, Tranfo G, Sapienza D, Scapellato ML, Bartolucci GB. Biological monitoring of low level exposure to benzene in an oil refinery: Effect of modulating factors. *Toxicology Letters*, 2018; 298, 70-75.
8. Keenan JJ, Gaffney SH, Galbraith DA, Beatty P, Paustenbach DJ. Gasoline: a complex chemical mixture, or a dangerous vehicle for benzene exposure? *Journal of Chemical and Biological Interaction*, 2010; 184(12): 293-295.
9. Badham HJ, LeBrun DP, Rutter A, Winn LM. Transplacental benzene exposure increases tumor incidence in mouse offspring: possible role of fetal benzene metabolism. *Carcinogenesis*. 2010; 31(6):1142-8.
10. Christian SG, Elekima I, Obisike UA, Aleru CP. Effect of petroleum on haematological parameters and lead level in fuel attendants in Port Harcourt, Nigeria. *International Journal of Science and Research (IJSR)*. 2016; 5(3): 280 -283
11. Frijhoff J, Winyard PG, Zarkovic N, Davies SS, Stocker R, Cheng D, Knight AR, Taylor EL, Oettrich J, Ruskovska T, Gasparovic AC, Cuadrado A, Weber D, Poulsen HE, Grune T, Schmidt HH, Ghezzi P. Clinical Relevance of Biomarkers of Oxidative Stress. *Antioxid Redox Signal*. 2015; 23(14): 1144-11470.
12. Ben-Chioma AE, Elekima I. Evaluation of Vitamin E and selenium levels in breast cancer patients in Port Harcourt Metropolis, Nigeria. *Journal of Advances in Medicine and Medical Research*. 2018; 28(2): 1-7.
13. George-Opuda MI, Adegoke OA, Bamigbowu EO. Effect of Vitamins C on Some Hematological Parameters in Albino Rats Treated with Gasoline. *Sciknow Publications*. 2014; 2(3): 55-58.
14. Friedewald WT, Levy RI, Fredrickson DS. Estimation of the concentration of low-density lipoprotein cholesterol in plasma without the use of the preparative ultracentrifuge. *Clinical Chemistry*, 1972; 18(6): 499–502
15. Uboh FE, Akpanabiatu EU, Eyoung EU, Ebong PE, Eka OU. Evaluation of Toxicological Implications of Inhalation Exposure to kerosene fumes and Petrol Fumes in Rats. *Acta Biology*. 2005; 49, 19-22.
16. Festus OO, Ebaluegbeifoh LO, Iyevhobu LOK, Dada FL, Iweka FK. Assessment of lipid profile in automobile mechanics in Ekpoma, Edo State. *European Journal of Biomedical and Pharmaceutical Sciences*. 2016; 3(9), 100-107
17. Ubani CS, Joshua PE, Umenwanne VC. Evaluation of toxicological implications of ingestion exposure to gasoline in mammals. *Biokemistri*. 2009; 21 (1): 33-39.
18. Chukwurah EF, Nnamdi AA, Chinedum CF, Ibe OE, Ogbodo S. Effects of inhalation of premium motor spirit fumes on plasma lipid profile and weight of albino rats. *Journal of Environmental Management and Safety*. 2020; 11(2): 1 – 11
19. Mohammad C, Ahmed Al-Sulivany BS, Fattah Y, Hadji MB. Effects of Gasoline and Smoking on Lipid Profile and Liver Functions among Gasoline Exposure Workers in Iraq. *International Journal of Occupational Safety and Health*. 2023; 13(1):19-28. 10.3126/ijosh.v13i1.43367
20. Ugbala JE, Adegoke OA, George-Opuda IM, Bamigbowu EO. Evaluation of lipid profile in male albino rats exposed to petrol fumes. *International Journal of Biomedical and Advance Research*. 2020; 11(2): 5321-5377.

21. **Ogbevire** L, Patrie O, Nathaniel M, John OD, Favour O, Ehis O, Richard O. Total Cholesterol, Triglyceride, High Density Lipoprotein Cholesterol and Low Density Cholesterol in Rat Exposed to Premium Motor Fumes. *North American Journal of Medical Science*, 2011; 3(6), 277-280.
22. **Kapil** S, Rakesh KS, Arun G, Vedpal Y. Impact of petroleum fumes on liver and kidney functioning of petrol filling attendants and garage attendants working in South Haryana, India. *European Journal of Pharmaceutical and Medical Research*. 2016; 3(8):569-573.
23. **Egbuonu** AC, Nkwazema DC, Ezeanyika L. (2015). Cardiovascular risks and impaired lipid metabolism in asymptomatic petroleum depot workers in Calabar metropolis. *Research Journal of Environmental Sciences*. 2015; 9 (6), 270-279
24. Gaur GS, Dixit AK. Comparative Study of Vitamin C on Serum Lipid Profile in Healthy Male and Female Human Subjects. *Journal of Scientific Research*. 2012; 4(3), 775–781.
25. Pehlivan FE. *Vitamin C: An Antioxidant Agent*. 2017; DOI: 10.5772/intechopen.69660
26. Sebastian JP, Arie K, Yaohui W, Peter Eck OK, Je-Hyuk L, Shenglin C, Christopher C, Anand D, Sudhir KD, Mark L. Vitamin C as an antioxidant: evaluation of its role in disease prevention. *Journal of America College of Nutrition*. 2003; 22(1):18-35. doi: 10.1080/07315724.2003.10719272.