Effects of ω-3 Fatty Acids on Liver Function Tests in Patients with Diabetes Mellitus

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Abstract

**Introduction:** Patients with type-2 Diabetes are augmented the risk of non alcoholic fatty liver disease that rises liver enzymes and total protein. To observe the favourable effects of fish oil capsule on liver function tests in patients with type 2 DM. **Materials and Methods:** A prospective interventional study was conducted from January 2017 to December 2017. A total numbers of 52 diagnosed type 2 diabetic patients of both sexes were scrutinized with age ranging from 40 to 50 years. Among them, 27 type 2 diabetic patients those who taken oral fish oil gel (2g/day) for 12 weeks were preferred as study group. Another 25 type 2 diabetic patients without supplementation of omega 3 fatty acid were nominated as control group for comparison. The study subjects were selected from Dhaka Medical College Hospital, Dhaka and personal contact from Dhaka city on the basis of criteria. The research work was administrated with ethical clearance from concerned authority.

**Results:** In this study ALP, LDH, total protein were reduced in diabetic patients after supplementation with omega-3 fatty acid in comparison to that of their baseline value. Again, after 12 weeks, ALP (Alkaline Phosphatase) LDH (Lactate dehydrogenase) and total protein were decreased in diabetic patients after supplementation with omega-3 fatty acid in comparison to control group. **Conclusion:** After analyzing the results of the study, it can be concluded that omega-3 fatty acid can reduce ALP, LDH and total protein levels in diabetic patients may be helpful to minimize the risk of fatty liver in type-2 diabetes mellitus.

**Key words:** Diabetes mellitus, Fatty liver, ALP, LDH, Total protein and Omega-3 fatty acid.

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**Introduction**

Diabetes mellitus (DM) is one of the multifactorial disorder characterized by hyperglycemia related with carbohydrate, protein and fat metabolism. The chronic hyperglycemia of diabetes is associated with long term damage of liver can causes non alcoholic fatty liver disease1. Diagnostic criteria of diabetes mellitus are fasting plasma glucose level ≥7.0mmol/l (126mg/dl) or plasma glucose 2 hours after an oral glucose ≥11.1 mmol /L (200mg/dl) and HbA1c≥6.5%. Within 2030, the extensiveness of diabetes mellitus will be 11.1 million in Bangladesh1. The prevalence of NASH is steadily increasing and prevalent in patients of Diabetes mellitus4.

The pancreatic hormone named insulin is needed for tissue development, growth and maintenance of whole body glucose homeostasis. Insulin maintains glucose homeostasis by improving the rate of glucose uptake into skeletal muscle and fat tissue. In the skeletal muscle, insulin fascilitates glucose uptake by stimulating translocation of GLUT-4 to plasma membrane3. Insulin resistance occurs when the insulin sensitive tissue loses response to insulin. The basic effect of insulin resistance on glucose metabolism is to oppose the uptake and utilization of glucose by most cells of the body. As a result blood glucose concentration rises, cell utilization of glucose falls, utilization of fat increases and free fatty acid level increases in blood6.

ω-3 Fatty acids are polyunsaturated fatty acids consists of alpha-linolenic acid (ALA), eicosapentaenoic acid (EPA) and docosahexanoic acid (DHA). They are found mainly in seafish including fatty fish (e.g. salmon, tuna and trout) and shellfish (e.g. crab, mussels and oysters). The omega 3 fatty acid stimulates insulin sensitivity, reduces blood clotting, improves fat digestion, boost up fertility, lessen depression and causes brain development in babys7.

Consumption of fish oil can decreases free fatty acid level, promotes insulin sensitivity as well as reduce the incidence of type 2 DM8. Omega-3 capsule act directly on insulin sensitive tissues, rises number of insulin receptors thus inhibits insulin resistance6. Intake of fish rich in omega-3 fatty acid, facilitate the action of insulin through various metabolic pathways, which are reduction of hepatic lipogenesis, suppression of the release of triglycerols from liver, improvement in ketogenesis, and oxidation of fatty acids in hepatic
Effects of ω-3 Fatty Acids on LFT in T2DM  

Habib, et al.

Effects of ω-3 Fatty Acids on LFT in T2DM

Habib, et al.

Cells 9. Non alchoholic fatty liver diseases are very common in type 2 diabetic patients 11. Moreover, intake of polyunsaturated fatty acid decreases serum ALP and GGT level 12.

Omega-3 fatty acid prevents this change by increasing peroxisome proliferator receptor gamma, increasing hepatic uptake and oxidation of free fatty acid in striated muscle 13. Therefore the present study is intended to assess the effect of supplementation of omega-3 fatty acid in Bangladesh diabetic patient that reduce liver enzymes.

Materials and Methods

This prospective, interventional study was administrated from Department of Physiology, Dhaka Medical College, Dhaka from January 2017 to December 2017. The research work was carried-out by obtaining ethical clearance from related departments, Research Review Committee and Ethical Review Committee of Dhaka medical college, Dhaka. The patients were selected from outdoor of Endocrinology, Dhaka medical college and personal contact from Dhaka city. At the beginning of study 60 diagnosed type-2 diabetic patients were randomly nominated on the basis of exclusion and inclusion criteria. There were 30 patients of control group and 30 patients of study groups scrutinized for study purpose. After 6 weeks of study period, 3 patients were relinquished from study group and 5 patients were dropped out from control group. Finally, total 52 type diabetic patient of both sexes with the age ranging from 40-50 years with FBG 7.0 mmol/l or 126 mg/dl, HbA1c 6.5%, serum total cholesterol 200 mg/dl, serum triglyceride > 150 mg/dl, LDL 130 mg/dl, BMI ≥ 30 Kg/m2 and patients with oral hypoglycemic drug were included in this study. Fatty liver was diagnosed by abdominal ultra-sonography subjects having history of heart, endocrine disorder, insulin therapy, viral hepatitis, acute or chronic infections, pregnant and lactating women were excluded from this study. For this study 27 diagnosed type-2 diabetic patients with omega-3 fatty acid supplementation were selected as study group and 25 type-2 diabetic patients without oral omega-3 fatty supplementation were selected as control group. The study group again segmented into pre-supplementation group and after 12 weeks of supplementation as post supplementation group. The control group was sub-divided as pre and post follow-up group. After selection, the nature, purpose and benefits of the study were explained to each subject and informed written consent was taken from participants. Before taking blood detailed family and medical history were taken. Anthropometric measurement of the subjects was recorded and blood pressure was measured. All the information were recorded in a prefixed questionnaire. With aseptic precaution, 5 ml of venous blood was obtained from ante-cubital vein by a disposable plastic syringe from each subject after overnight fasting for biochemical tests. Serum ALP and GGT was estimated by enzymatic colorimetric method in auto-analyzer in department of Laboratory Medicine Dhaka Medical College Hospital, Dhaka. Omega-3 fatty acid (2gm) was supplied to study group then they were asked to intake twice daily for 12 weeks with proper directions. Patients were instructed not to change their diet and physical activities during the course of the study. A regular telephonic communication and periodic visit was made to participants because most of them are staff of Dhaka medical college. For statistical analysis, Paired Student’s ‘t’ test and Unpaired Student’s ‘t’ test were performed as applicable using SPSS for windows version 16.0. Data were expressed as mean ± SE. The p value of < 0.05 was accepted as level of significance.

Results

In this study no significant difference were seen in age, sex, BMI, systolic and diastolic blood pressure between study and control group (Table I).

Table I: General characteristics of the patients in both groups (N=52).

| Parameters                  | Study group (n=27) | Control group (n=25) |
|-----------------------------|-------------------|----------------------|
| Age (years)                 | 45.90 ± 3.80      | 44.92 ± 3.75         |
| Sex (%)                     |                   |                      |
| Male                        | 18 (66.7%)        | 11 (44%)             |
| Female                      | 9 (33.3%)         | 14 (56%)             |
| BMI (kg/m²)                 | 25.03 ± 2.27      | 25.87 ± 1.75         |
| Systolic BP * (mmHg)        | 119.07 ± 7.08     | 121.79 ± 4.47        |
| Diastolic BP * (mmHg)       | 79.63 ± 6.26      | 80.00 ± 0.00         |
| Duration of disease * (years)| 5.43 ± 1.50       | 5.35 ± 1.57          |

Results were expressed as mean ± SD. a=Unpaired Student’s ‘t’ test was performed to compare between the groups. b= Chi Square test was performed to compare male and female between the groups. The test of significance was calculated and p value < 0.05 was accepted as level of significance. N= total number of subjects, n = number of subjects in each group ns= non-significant */**/***= significant. T2DMs=Type 2 diabetes mellitus with supplementation T2DM=Type 2 diabetes mellitus without supplementation.

In this study, the mean serum ALP and GGT levels were almost similar and there is no statistical difference were observed at the beginning of the study. In study group, the mean serum ALP (p<.01) and mean serum GGT (p<.01) level were found remarkable lowe in post supplementation group, than pre-supplementation group. Again the mean serum ALP (p<.01) levels and mean serum GGT (p<.01) levels were found significantly lower and in study group compared to control group. In control group, there was no statistical dissimilarity were perceived in mean serum ALP and GGT between pre-follow-up and post follow-up group.
Effects of ω-3 Fatty Acids on LFT in T2DM

Rebecca, et al.

Reduction in blood. The consequence of in skeletal muscle. Consequently, free fatty acid level is free fatty acid. It also uprise the free fatty acids oxidation acid. An increase in PPAR–α leads to hepatic uptake of which increase in number in presence of omega-3 fatty acid. Peroxisome proliferator receptor–α exists in the liver acid has a role on reducing serum triglyceride level. Omega-3 fatty acid are influenced by rising serum TG level. Omega-3 fatty acids has a role on reducing serum triglyceride level. Peroxisome proliferator receptor–α exists in the liver which increase in number in presence of omega-3 fatty acid. An increase in PPAR–α leads to hepatic uptake of free fatty acid. It also uprise the free fatty acids oxidation in skeletal muscle. As a result, free fatty acid level is reduced in blood. The consequence of free fatty acid reduction helps to decrease triglyceride synthesis. Thus, fish oil capsules reduces serum triglyceride level that promotes the binding of insulin to its receptor and enhances insulin sensitivity. Fish oil activate PPAR–α (peroxisome proliferator activated receptor alpha) and down regulate sterol regulatory element binding protein-1c (SREBP-1) that enhance fatty acid oxidation and decreases liver enzymes. Modulation of Alkaline Phosphatase enzyme can be done by dietary fish oil. Fish oil supplements lowers plasma lactate dehydrogenase that’s releases from endoplasmic reticulum and improve glucose tolerance by molecular mechanism. In diabetic patient total protein level rises. Supplementation of fish oil capsule decreases protein synthesis signaling so total protein decreases.

In the present study serum ALP, LDH and Total protein levels decreases in patients with T2DM after supplementation of omega-3 fatty acid in contrast to their baseline value and control group. Omega-3 fatty acid supplementation lessen ALP and LDH by increasing fatty acid oxidation and alter signaling pathway of protein and decreases total protein. This premises the binding of insulin to its receptor and improves insulin sensitivity.

Discussion

In the present study, the mean serum ALP and GGT levels were reduced in patients of T2DM after supplementation with omega-3 fatty acid in comparison to that of their baseline value. Again, after 12 weeks, mean high density lipoprotein (HDL) level was significantly higher in type-2 diabetic patients supplemented with omega-3 fatty acid in comparison to that of diabetic control group without omega-3 fatty acid capsule. Insulin resistance leads impaired fatty acid oxidation in liver that causes accumulation of fat in liver. When diabetic patient suffers from non alcholohic fatty liver diseases ALP and GGT rises. Almost similar type of result were seen by different researchers of different countries. On the contrary, found no significant difference in lipid profile in patients after supplementation of omega-3 fatty acid in comparison to that of their baseline values and diabetic control group who were not supplemented with omega-3 fatty acid. There were a history of less physical activity in control group who were not supplemented with omega-3 fatty acid in comparison to that of their baseline values and diabetic patients after supplementation of omega-3 fatty acid in contrast to their baseline value and control group. Omega-3 fatty acid supplementation lessen ALP and LDH by increasing fatty acid oxidation and alter signaling pathway of protein and decreases total protein. This premises the binding of insulin to its receptor and improves insulin sensitivity.

Conclusion

After analyzing the results of the study, it can be concluded that supplementation of omega-3 fatty acid can reduces serum ALP and GGT levels in patients with type-2 diabetes mellitus. Therefore, omega-3 fatty acid containing diet may be useful to keep down the complications in type-2 diabetes mellitus.

Conflict of Interest: None.

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Table-II: Serum Alkaline phosphatase, serum lactate dehydrogenate and total protein levels in different groups (N=52).

| Parameters          | Study group (n=27) | Control group (n=25) |
|---------------------|--------------------|----------------------|
|                     | Pre-supplementation | Post supplementation | Pre-supplementation | Post supplementation |
| ALP (IU/L)          | 74.6 ± 2.71        | 38.5 ± 1.67          | 69.20 ± 4.20        | 61.96 ± 4.43          |
| LDH (IU/L)          | 150.3 ± 4.03       | 142.5 ± 1.02         | 147.00 ± 2.64       | 146.6 ± 2.21          |
| Total protein (g/dl)| 11.06 ± 2.01       | 8.1 ± 1.1            | 10.06 ± 1.02        | 9.8 ± 1.4             |

Results are expressed as mean ± SD. a= Paired student’s t test was performed for comparison within groups and b=unpaired t test was performed to compare between groups. p value < 0.05 was accepted as level of significance. N= total number of subjects, n = number of subjects in each group, ALP=Alkaline Phosphatase, GGT=Gamma glutamyl transeferase (*= study group baseline vs study group after 12 weeks of supplementation; # = study group after 12 weeks vs control group after 12 weeks); (* p<.01, **p<0.001;# p<.01,## p<0.001).
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