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The Effect Of Eicosapentanoic Acid Administration On Clinical Outcome Of Patients With Major Depression: A Pilot Study

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ABSTRACT

Background: Major depressive disorder (MDD) is a common brain disorder that affects approximately 10% of the world population and leads to significant disability. The current study was aimed to evaluate the impact of Omega3 PUFAs administration on the clinical outcome of patients with depression.

Patients and Methods: This prospective, randomized controlled study included a total of Forty-two patients who diagnosed with depression according to a strict inclusion and exclusion criteria, attending at the outpatient clinics of the Department of Psychiatry, Al-Zahraa University Hospital, Cairo, Egypt. Approval of the Ethical Research Committee of Faculty of Pharmacy, Ain Shams University (registration number 61), and a written informed consent from all the subjects were obtained. This study was conducted between February 2015 till August 2016. patients were randomly assigned to either; Group 1; (intervention n=21); received the prescribed antidepressant + omega3 (2100mg) for 8 weeks, or Group 2; (control, n=21); received the prescribed antidepressant only for 8 weeks. Baseline evaluation and 8-week assessment included; patient demographic-data collection, history taking and clinical assessment of DSM-5 criteria & HAM-D score. Laboratory assessment included; complete blood picture (CBC), prothrombin time (PT), activated partial thromboplastin time (aPTT). Patients were followed up regularly every 2 weeks for 8 weeks for the occurrence of side effects due to antidepressants/ Omega 3 and compliance with medications.

Results: The 2 groups were comparable at baseline. The test group showed a significant improvement in the HAMD score from baseline values and versus the control. There was no significant difference in the reported side effects between the 2 groups.

Conclusion: Omega -3 PUFAs administration at a dose of 2100 (EPA1350 mg/ , DHA 600 mg/ 150 mg other omega 3 FA) for 8 weeks, improved depression symptoms and was well tolerated.

Keywords: Omega-3 PUFAs, depression, HAMD, EPA/DHA.

INTRODUCTION

Major depressive disorder (MDD) is a common brain disorder that affects approximately 10% of the world population. According to the Diagnostic and Statistical Manual of Mental Disorder, Fifth Edition (DSM-5, 2013), MDD is characterized by the loss of interest in pleasure, low self-esteem, disturbed sleep or appetite, fatigue, and diminished ability to think or concentrate. These problems often become chronic and recurrent, and at the worst, can lead to suicide (1).

Major depression is a commonly occurring, serious, recurrent disorder linked to diminished role functioning and quality of life, medical morbidity, and mortality. The World Health Organization (WHO) has ranked depression as the fourth leading cause of disability worldwide and projects that, by 2020, it will be the second leading cause (2).

The lifetime prevalence of depression in urban and rural Egyptian populations found to be 11.4 and 19.7% (3).

New-generation antidepressants appear more effective than older drugs in treatment of major depression, with response rates of up to 50%, but they do not effectively treat all depressed patients. In addition, many drugs have side effects that can affect compliance and morbidity. So patients are increasingly using complementary and alternative medicine (CAM) therapies to treat depression (4). One such possibility is the use of n-3 omega polyunsaturated fatty acids (PUFAs) for the treatment of depression. A link between omega-3 fatty acids and mood disorders has been suggested by some studies showing a lower incidence of depression among populations with a diet rich in omega-3 fatty acids (5).

Omega-3 PUFA have been proposed, for treatment of Major depressive disorder. The positive effects of omega-3 PUFA on depression may depend on their physiological abundant content in the human nervous system and their involvement in neurogenesis and neuroplasticity (6). Moreover, their anti-inflammatory capacity may counteract inflammatory processes occurring in depression (7). Also lower concentrations of omega 3 PUFA in plasma or red blood cells may contribute to depression. Administration of omega 3 PUFAs significantly improved depressive symptoms in patients with major depression (8).
Several studies used omega-3 polyunsaturated fatty acids (PUFA) eicosapentaeoic acid (EPA) and docosahexanoic acid (DHA) in similar doses with varying results.

It has been previously documented that EPA at ratios > 60% can positively affect depression outcome. EPA and DHA are structurally similar and might be expected to compete in approximately a 1:1 ratio for binding sites. Thus the amount of EPA unopposed by DHA may be critical for effective PUFA supplementation in treatment of depressive episodes (9).

Hence the aim of the current study was to evaluate the impact of Omega-3 PUFAs administration (using an EPA in a dose more than twice the concentration of DHA), on the clinical outcome of patients with depression.

PATIENTS AND METHODS

This prospective, randomized controlled study included a total of Forty-two patients who diagnosed with depression according to a strict inclusion and exclusion criteria, attending at the outpatient clinics of the Department of Psychiatry, Al-Zahraa University Hospital, Cairo, Egypt. Approval of the Ethical Research Committee of Faculty of Pharmacy, Ain Shams University (registration number 61), and a written informed consent from all the subjects were obtained. This study was conducted between February 2015 till August 2016. Patients were randomly distributed into Group1; (intervention n=21); received the prescribed antidepressant + omega3 (2100mg) for 8 weeks, and Group 2; (control, n=21); received the prescribed antidepressant only for 8 weeks.

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Patient demographic-data collection, history taking and clinical assessment including, assessment of DSM-5 criteria by psychiatric interview and clinical pharmacist assessment of HAM-D score to assess the severity of depression.

Laboratory assessment including: complete blood picture (CBC), prothrombin time (PT), activated partial thromboplastin time (aPTT). Patients were followed up regularly every 2 weeks for 8 weeks for the occurrence of side effects due to antidepressants/ Omega 3 and compliance with medications.

Inclusion criteria included: an age range of (18-65 years); a diagnosis of depression according to the Diagnostic statistical manual for mental disorders 5th Edition DSM-5; and a Hamilton Depression Rating Scale (HAM-D) score of greater than or equal to 17. Exclusion criteria included having any of the following, a psychotic or seizure disorder; a current drug/ alcohol abuse or dependence; or history of drug or alcohol abuse or dependence within the previous six months; an unstable medical or neurological condition that is likely to interfere with the treatment of depression; pregnancy; history of allergy to omega-3 fatty acids, or shellfish; concomitant therapy with psychotropic medications; active suicidal ideation or other safety concerns; exposed to treatment with fluoxetine or monoamine oxidase inhibitors (MAOIs) in the previous two months; on anticoagulant therapy; receiving dietary intake of omega-3 at baseline; or refusal to participate in the study.

All procedures performed in studies involving human participants were in accordance with the ethical standards of the Institutional and/or National Research Committee and with the 1964 Helsinki declaration and its later amendments or comparable ethical standards.

Forty eligible patients were randomly assigned to either:

Group1; (intervention n=21); received the prescribed antidepressant + Ocean blue professional omega3 2100 (Eicosapentanoic acid 1350 mg, Decosahexanoic acid 600 mg and 150 mg other omega 3 fatty acids) for 8 weeks.

Group 2; (control, n=21); received the prescribed antidepressant only for 8 weeks.
CBC, PT, aPTT, and depression outcome using HAM-D score.

The occurrence of side effects to medications was assessed using patient side-effect reporting cards that patients had to fill daily in between visits.

HAM-D score was interpreted using the following scores; 0-7 indicated no depression; 8-16, mild depression; 17 – 23, moderate depression, and > 23, indicated severe depression (10).

A response to treatment was defined as at least a 50% reduction in the HAM-D score, while “Remission” is a return to a normal state or a HAM-D of 7 or less.

The Primary end point was comparing the differences in HAMD score from base line to 8 weeks between groups. Other end points were comparing the difference in side effect occurrence between groups.

Statistical Analysis

Data management and analysis was performed using Statistical Package for Social Sciences (SPSS) vs. 23. Numerical data was summarized using means and standard deviations or medians and ranges, as appropriate. Categorical data was summarized as numbers and percentages. Numerical data was explored for normality using Kolmogrov-Smirnov test and Shapiro-Wilk test. Exploration of data revealed that the collected values were not normally distributed. Comparisons between both groups was done by Mann-Whitney test. Change between baseline and end of study was tested using the Wilcoxon Rank Signed. All p-values were two-sided. P-values < 0.05 were considered significant.

RESULTS

From February 2015 till August 2016, out of a total of 75 depression patients screened, only 42 fulfilled the inclusion criteria & were included in the study.

Baseline laboratory evaluation and patients’ demographics were comparable between the 2 groups as showed by table 1.

Table (1): Comparison between control and intervention groups in demographics, laboratory and clinical parameters.

<table>
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<th>Group</th>
<th>P-values</th>
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After 8 weeks, the intervention group showed a significant decline in the overall HAMD score and the percentage change over time versus the control as showed by figure 1.

Figure (1): Box plot of HAMD score over time between control and intervention groups.

Over the study period, there was no significant difference between the 2 groups in WBCs, Hemoglobin or platelets counts. The intervention group showed a significant difference in the percentage change over time of the aPTT versus the control.

There was no significant difference between the 2 groups in the occurrence of side effects, namely; sexual dysfunction, dizziness, GIT upset or sleep abnormalities.
The side effects reported in the intervention group were non-remakable except for; fishy- after taste was reported in 8 patients (38.1%) , itching was reported in 4 patients (19%) and GIT upset was reported in 4 patients (19%). All of these side effects were mild and self-limited and did not require drug discontinuation.

DISCUSSION

Major depressive disorder (MDD) is the most prevalent Axis I disorder and one of the leading causes of disability in the United States and worldwide. In the global burden of disease (GBD) 2000, depressive disorders were ranked the third leading cause of burden (equivalent to 4.3% of all disability adjusted life years (DALYs)) after lower respiratory infections and diarrheal diseases (11).

The lifetime prevalence of depression in urban and rural Egyptian populations was reported to be 11.4 and 19.7% (3).

Pharmacotherapy is the mainstay of treatment for depression, yet an adequate response to a single antidepressant medication, measured as an attenuation of 50% or more in the severity of depressive symptoms, only occurs in about 50%-75% of patients. Moreover, remission rates are even less and do not usually exceed 30% with any single agent. Hence, the search for a naturally occurring, inexpensive, safe, and effective agent continues. The use of n-3 omega polyunsaturated fatty acids (PUFAs) for treatment of depression is a potential option (5).

The results of the current study showed that the addition of omega 3 fatty acids to the standard antidepressant treatment was associated with improvement of both HAM-D score and depression symptoms.

The current study used 2100 mg omega 3 fatty acids, which is comprised of 1350 mg eicosapentanoic acid, 600 mg docosahexanoic acid and 150 mg decosapentanoic acid in addition to the traditional selective serotonin reuptake inhibitors used for depression. The rationale for this high concentration of EPA is the previous evidence reporting that EPA at doses more than twice the dose of DHA can positively affect depression outcome (9).

Moreover it has been shown that EPA and DHA are structurally similar and might be expected to compete in approximately a 1:1 ratio for binding sites. This explanation implies a functional competition not only between omega-3 and omega-6 PUFAs but also within omega-3 species, with regard to depression. Thus EPA in excess of DHA may be considered mechanistically to be unopposed EPA and so will be considered as the active component of PUFAs which will exhibit a therapeutic outcome with regard to depression treatment (9).

The Hamilton Rating Scale for Depression (HAM-D) used in this study, is a clinician rating scale, which is used to assess the severity of major depression and to assess the response to treatment. It includes 21 items that the clinician has to ask the patient about. To assess disease severity, a score from; 17-19 is moderate depression, 20-22 is severe depression, > 22 is very severe depression (10).

In the current study, baseline HAM-D score was comparable in both groups. The omega 3 fatty acids group showed a significant improvement in HAM-D score levels as compared to baseline values and to the control.

Similarly several studies have supported this finding. M H Rapaport et al, evaluated whether inflammatory biomarkers act as moderators of clinical response to omega-3 (n-3) fatty acids in subjects with major depressive disorder (MDD). One hundred fifty-five subjects with (DSM-IV) MDD, with a baseline (HAM-D-17) score of 15 and baseline biomarker data (interleutkin (IL)-1ra, IL-6, high-sensitivity C-reactive protein (hs-CRP), leptin and adiponectin) were randomized for about 8 weeks to double-blind treatment with eicosapentanoic acid (EPA) 1060 mg, docosahexanoic acid (DHA) 900 mg or placebo. Outcomes were determined using mixed model repeated measures analysis for ‘high’ and ‘low’ inflammation groups based on individual and combined biomarkers. Results showed significant changes in HAM-D-17 from baseline to treatment week 8 between intervention and placebo group (12).

Gertsik, L et al evaluated 42 subjects with MDD in a nine week randomized, masked, placebo-controlled study of combination therapy (2 grams containing a blend of 900mg EPA, 200mg DHA, and 100mg other omega-3 fatty acids twice daily + citalopram) versus monotherapy (2 grams olive oil per day plus citalopram) treatment of MDD. The combination therapy demonstrated significantly greater improvement in HAMD scores beginning at week 4 and was more effective than monotherapy in decreasing signs and symptoms of MDD during the 8 weeks of active treatment (5).
Su et al, conducted an 8-week, double-blind, placebo-controlled trial, comparing omega-3 PUFAs (6.6 g/day) with placebo, on the top of the usual treatment, in 28 patients with major depressive disorder. Patients in the omega-3 PUFA group had a significantly lower score on the HAMD score than those in the placebo group. In another study by Su et al on Perinatal depression, an 8-week, double-blind, placebo-controlled trial comparing omega-3 PUFAs (3.4 g/d) with placebo in pregnant women with (DSM-IV criteria) was conducted. No psychotropic agent was given 1 month prior to or during the study period. The results showed that subjects in the omega-3 group had significantly lower HAM-D scores at weeks 6 and 8, a significantly higher response rate (62% vs. 27%, \( p = .03 \)), and a higher remission rate. At the study end point, subjects in the omega-3 group also had significantly lower depressive symptoms ratings on the used scores and no adverse effects on the subjects and newborns were reported from Omega 3 administration.

Frangou and colleagues randomized 75 depressed subjects in a double-blind trial to receive 1 g/d ethyl-EPA, 2g/d ethyl-EPA, or placebo for 12 weeks. EPA outperformed placebo significantly at both ethyl-EPA doses, based on HAM-D scores; the higher dose of ethyl-EPA seemed to confer no added benefit in comparison with 1 g/d.

Peet and Horrobin conducted a randomized, placebo-controlled, dose-finding study of ethyl-eicosapentaenoate (ethyl-EPA) as adjunctive therapy for 70 adults with unipolar major depressive disorder (MDD) who were not responsive to standard antidepressants. A dose of 1 g/d ethyl-EPA for 12 weeks yielded significantly higher response rates (53%) than placebo (29%), with notable improvement of depressed mood, anxiety, sleep disturbance, libido, and suicidality. The 2 g/d group showed no significant difference between drug and placebo, and the 4 g/d group showed a non-significant trend toward improvement. These results suggested a therapeutic window for n-3FA required for maximum benefit, and it is possible that an “overcorrection” of the n-6FA:n-3FA ratio with higher n-3FA doses may limit the antidepressant effect of ethyl-EPA.

In contrast, Careny et al, evaluated 122 patients with major depression and CHD who were randomized in double-blind fashion to receive 50 mg/d of sertraline and 2 g/d of omega-3 acid ethyl esters (930 mg of EPA and 750 mg of DHA) or to corn oil placebo capsules for 10 weeks. The results of this study showed omega-3 fatty acids did not result in superior depression symptoms reduction (by BDI-II and HAM-D) at 10 weeks, compared with sertraline and placebo. The discrepancy in these results with others could be attributed to the presence of CHD with depression and the smaller ratio between EPA/ EPA which was not twice the DHA.

Moreover in a study by Silvers et al, 77 participants were randomly assigned to receive 8 g of fish or olive oil per day in addition to their existing therapy. Fish oil was no more effective than the control as an add-on therapy for depression in this setting. These results could be attributed to the higher omega3 dose of 8g/day, as doses ≥ 4g were proven to be non-effective and attained a non-linear dose effect. “Overcorrection” of the n-6FA:n-3FA ratio with higher n-3FA doses may limit the antidepressant effect of omega 3 fatty acids.

The omega-3 fatty acids have been shown to be very safe and well tolerated. The most reported side effects, such as gastrointestinal upset and fishy aftertaste, tend to occur with higher doses (greater than 5 g/d) and with less pure preparations. At the more typical doses of 1 g/d with highly purified omega-3 preparations, these adverse effects are less common. There has been some concern about the possibility of bleeding with doses greater than 3 g/d, although this risk seems to be minimal, except in patients who are taking other agents that also affect platelet function.

The current study did not report any significant side effects with Omega-3 that required intervention or therapy discontinuation.

Similarly, Kiecolt-Glaser and colleagues reported non-serious adverse events with omega 3 fatty acids administration including, sore throat, nasal symptoms, stomach pain, tachycardia, and Diarrhea with non significant differences between intervention and control group.

In a review by Kris-Etherton et al, it was reported that the FDA has ruled that intakes of up to 3 g/d of marine omega-3 fatty acids are GRAS (Generally Recognized As Safe) for inclusion in the diet. Perhaps the most common side effect was a fishy aftertaste.

In a 6-month trial providing 275 patients with 6.9 g of EPA+DHA in 10 capsules daily, there was no difference between the fish oil and corn oil.
control groups for any adverse event. GIT upset was reported by 8% of the latter and 7% of the former. Omega 3 preparations appear to be well tolerated with non serious adverse events [21].

CONCLUSION

Omega 3 PUFAs administration resulted in improvement of major depressive symptoms indicated by a HAMD score reduction. Omega 3 PUFA was well tolerated and patients who experienced side effects did not require any drug-intervention, dose reduction or dose discontinuation. Omega-3 PUFAs at a dose of 2100 (Eicosapentanoic acid 1350 mg, Decosahexanoic acid 600 mg and 150 mg other omega 3 fatty acids) for 8 weeks is a well tolerated supplement that improves depressive symptoms & reduces HAMD scores. Conflict of Interest: The authors declare that they have no conflict of interest. DISCLAIMERS: The use of any specific brand of neurotrophic factor was not intentional in the trial. Any available type of neurotrophic factor was used regardless of the producing company.

REFERENCES


