



Salud mental

ISSN: 0185-3325

Instituto Nacional de Psiquiatría Ramón de la Fuente Muñiz

Álvarez-Ramírez, Miriam; Lara, Ma. Asunción; Cervantes-Alfaro, José Miguel; Robinson-Fuentes, Virginia Angélica; Alveano-Hernández, Jesús

Correlation between intake of omega-3 fatty acids and depressive and anxiety symptoms during pregnancy

Salud mental, vol. 41, no. 1, 2018, January-February, pp. 31-38

Instituto Nacional de Psiquiatría Ramón de la Fuente Muñiz

DOI: 10.17711/SM.0185-3325.2018.002

Available in: <http://www.redalyc.org/articulo.oa?id=58255540005>

- How to cite
- Complete issue
- More information about this article
- Journal's webpage in redalyc.org

UDEM  redalyc.org

Scientific Information System Redalyc

Network of Scientific Journals from Latin America and the Caribbean, Spain and Portugal

Project academic non-profit, developed under the open access initiative

Correlation between intake of omega-3 fatty acids and depressive and anxiety symptoms during pregnancy

Miriam Álvarez-Ramírez,¹ Ma. Asunción Lara,² José Miguel Cervantes-Alfaro,³ Virginia Angélica Robinson-Fuentes,¹ Jesús Alveano-Hernández⁴

¹ División de Estudios de Posgrado, Laboratorio de Química Analítica, Facultad de Medicina Dr. Ignacio Chávez, Universidad Michoacana de San Nicolás de Hidalgo, Morelia, Michoacán, México.

² Departamento de Modelos de Intervención, Instituto Nacional de Psiquiatría Ramón de la Fuente Muñiz, Ciudad de México, México.

³ División de Estudios de Posgrado, Laboratorio de Neurociencias y Neuroprotectores Endógenos, Facultad de Medicina Dr. Ignacio Chávez, Universidad Michoacana de San Nicolás de Hidalgo, Morelia, Michoacán, México.

⁴ Escuela de Medicina, Universidad Vasco de Quiroga, Morelia, Michoacán, México.

Correspondence:

Miriam Álvarez Ramírez
División de Estudios de Posgrado,
Laboratorio de Química Analítica,
Facultad de Medicina Dr. Ignacio
Chávez, Universidad Michoacana de
San Nicolás de Hidalgo.
Astrónomos No. 28, Fracc Cosmos.
C.P. 58050, Morelia, Michoacán,
México.
Phone: (52) 443 155-8067
Email: malvarez.nutricion@gmail.com

Received: 14 March 2017

Accepted: 25 January 2018

Citation:

Álvarez-Ramírez, M., Lara, M. A., Cervantes-Alfaro, J. M., Robinson-Fuentes, V. A., Alveano-Hernández, J. Correlation between intake of omega-3 fatty acids and depressive and anxiety symptoms during pregnancy. *Salud Mental*, 41(1), 31-38. doi: 10.17711/SM.0185-3325.2018.002



ABSTRACT

Introduction. Anxiety and depression during pregnancy are disabling disorders associated to complications during the pregnancy, delivery, and postpartum period, with a significant prevalence, between 9 and 20%, hence its importance. Nutritional factors, such as omega-3 fatty acids (ω -3FA) deficiency, have been related with both disorders during pregnancy, especially docosahexaenoic acid (DHA) and eicosapentaenoic acid (EPA). **Objective.** To evaluate the association between dietary intake of EPA and DHA, and anxiety symptoms (AS) and depressive symptoms (DS) in Mexican pregnant women. **Method.** The sample consisted of 151 women in the second trimester of pregnancy. Instruments included a Food Frequency Questionnaire, the Trait Anxiety Inventory, and the Edinburgh Postnatal Depression Scale. **Results.** A daily intake of DHA and EPA of 70 mg/d and 30 mg/d, respectively, was found. The prevalence of AS was 44.4%, and 17.9% for DS. We also found the correlation negatively moderate between dietary intake of DHA and EPA and AS ($p = .003$, $p = .017$) and DS ($p = .001$, $p = .020$) in the group of women who had a severely insufficient intake of ω -3FA. **Discussion and conclusion.** The statistical significance shows a negative correlation between variables and the dietary intake of ω -3FA explains in a very small percentage the variability of AS and DS, according to their coefficient of determination. These results suggest the need for an investigation of this relationship through interventional studies.

Keywords: Fatty acids, omega-3, prenatal anxiety, prenatal depression.

RESUMEN

Introducción. La prevalencia de ansiedad y depresión en el embarazo oscila entre 9 y 20%. Poseen una gran importancia al ser trastornos incapacitantes asociados a complicaciones durante el embarazo, parto y postparto. Factores nutricionales, como la deficiencia de ácidos grasos omega-3 (AG ω -3), se han relacionado con ambos trastornos durante el embarazo, especialmente el ácido docosahexaenoico (DHA) y el ácido eicosapentaenoico (EPA). **Objetivo.** Evaluar la asociación entre la ingesta dietética de EPA y DHA y los síntomas de ansiedad (SA) y depresión (DS) en mujeres embarazadas mexicanas. **Método.** Se entrevistó a 151 gestantes fueron entrevistadas en el segundo trimestre de embarazo. Los instrumentos de medición utilizados fueron: un Cuestionario de Frecuencia de Consumo de Alimentos, el Inventario de Ansiedad-Rasgo y la Escala de Depresión Postnatal de Edimburgo. **Resultados.** Se encontró una correlación negativa moderada entre la ingesta de DHA y EPA y los SA ($p = .003$, $p = .017$) y SD ($p = .001$, $p = .020$) en el grupo de mujeres con ingesta gravemente insuficiente. Se encontró una ingesta dietética diaria de DHA y EPA de 70 mg y 30 mg, respectivamente. La prevalencia de SA fue 44.4% y de SD, 17.9%. **Discusión y conclusión.** Aunque la significancia estadística mostró una correlación negativa entre variables, de acuerdo con el coeficiente de determinación, la ingesta dietética de AG ω -3 explica en un escaso porcentaje la variabilidad de los SA y SD. Estos resultados sugieren la necesidad de investigar dicha relación por medio de estudios de intervención.

Palabras clave: Ácidos grasos, omega-3, ansiedad prenatal, depresión prenatal.

INTRODUCTION

Pregnancy is a biopsychosocial condition that can increase the risk of mental disorders such as anxiety and depression in nearly 15.6% of pregnant women (Rahman et al., 2013). Anxiety symptoms (AS) during pregnancy have an international prevalence between 9.1% and 15.6% (Fadzil et al., 2013; Rubertsson et al., 2014) and a prevalence between 10% and 15% in Mexico (Ceballos-Martínez et al., 2010; Navarrete et al., 2012). Occasionally, AS during pregnancy are more frequent than depressive symptoms (DS) (Navarrete et al., 2012). A recent Mexican study shows a 16.6% prevalence of DS during pregnancy (Lara et al., 2015). AS and DS are not isolated disorders, they co-occur in 30% to 58% of patients with major depressive disorder (Field et al., 2010).

Both AS and DS have a negative impact in women's lives, their children, and their families. The World Health Organization recommends promoting low-cost interventions to improve maternal mental health (de Castro, Place, Villalobos, & Allen-Leigh, 2015; Berenzon, Lara, Robles, & Medina-Mora, 2013). However, these disorders do not represent a priority for the health care systems in Mexico, despite being associated to important effects on mother, fetus, neonate, and child (Ding et al., 2014; Hanley & Oberlander, 2014). Depressed women have difficulties at work, higher prevalence of cardiovascular disease, stroke, and type 2 diabetes. Also, depressed and anxious mothers neglect their health and the health of their future child, leading to a deficient mother-baby bonding (de Castro et al., 2015). Furthermore, the fetuses of prenatally depressed women also show less total movement and an increased heart rate, have a delayed fetal growth, low fetal and birth weight, and low fetal length, with their subsequent risk of disease in adult life. Besides, newborns of depressed mothers are less responsive to stimulation in addition to other effects (Field, 2011).

There are many psychosocial risk factors which play an important role in the etiology of AS and DS. In Mexico, SD prevail in socioeconomic vulnerability and marginalization contexts (Lara, 2014; de Castro et al., 2015).

Recently, nutritional factors have been associated with AS and DS. The most studied nutritional risk factor is the intake of omega-3 polyunsaturated fatty acids (ω -3FA), especially their long-chain derivatives: docosahexaenoic acid (DHA 22:6 ω -3) and eicosapentaenoic acid (EPA 20:5 ω -3). DHA and EPA are endogenously synthesized from α -linolenic acid (ALA 18:3 ω -3), but in a very limited amount, 9% is converted to DHA and 21% to EPA (Assies et al., 2010). Therefore, their intake should come from the diet, that is, from sources such as tuna, salmon, sardines, trout, school shark, and seafood. The current adequate intake, issued by the Institute of Medicine of the National Academies (IOM), for DHA, EPA, and ALA during pregnancy and lactation, are 300 mg/d, 220 mg/d and 1.4 g/d, respectively (Simopou-

los, 2011). During the last trimester of pregnancy 67 mg/d of DHA are transferred to the fetus for the growth and development of its brain and retina. About 50% to 60% of the dry weight of an adult brain is composed of lipids, of which 20% corresponds to DHA (Liu, Green, Mann, Rapoport, & Sublette, 2015). Thus, if the intake of ω -3FA by the mother is insufficient, it is common that pregnancy depletes the mother's stock of cerebral and erythrocyte ω -3FA (Morse, 2012). This deficiency, in the mother, is associated with a higher risk of mood disorders and preeclampsia, for example, whereas, in the newborn, it is associated with low neural development index, preterm birth, low height, and weight at birth (Van Eijsden, Hornstra, van der Wal, Vrijkotte, & Bonsel, 2008).

An inverse association between a per capita intake of fish and an annual prevalence of depression was observed in the last 15 years (Grosso et al., 2014), which led to propose that dietary deficiency of ω -3FA (measured as plasmatic, erythrocyte, and brain concentrations) is associated with higher depression, both in the perinatal period (Hibbeln, 2002; Markhus et al., 2013) and in other stages of life (Hoffmire, Block, Thevenet-Morrison, & van Wijngaarden, 2012; Grosso et al., 2014; Su, Matsuoka, & Pae, 2015). There is a relation between the intake of ω -3FA and prenatal AS in animal and human models (Mizunoya et al., 2013; Vaz et al., 2013; Liu et al., 2015; Su et al., 2015).

The biological mechanisms proposed to explain this relation can be grouped according to the effect of ω -3FA in:

- Functionality of the neuronal plasmatic membrane. DHA confers greater fluidity and permeability to plasma membrane, allowing an appropriate physical environment for the integral activity of different components present in neuronal membrane (Bazan, Musto, & Knott, 2011).
- Neuroendocrine and neuroinflammatory modulation. EPA and DHA are precursors of eicosanoids and docosanoids, biologically active lipid mediators, which have a wide variety of anti-inflammatory activities related to their antidepressant effect (Grosso et al., 2014).
- Neurotransmission. DHA participates in serotonergic, dopaminergic, and endocannabinoid neurotransmission systems. Its deficiency can affect the metabolism, release, recapture of neurotransmitter, and functionality of its receptors (Rizzo et al., 2012).
- Synaptic plasticity. DHA participates in the synthesis of synaptic plasticity proteins in the hippocampus, like the brain-derived neurotrophic factor, which exerts antidepressant effects (Grosso et al., 2014).

Pregnancy is an adequate physiological condition to study the relation of ω -3FA and mental disorders for the ease with which maternal cerebral supplies of DHA are depleted (Markhus et al., 2013), and also because of the vulnerability of pregnant women to AS and/or DS (Wisner, Lara-Cinisomo, Pinheiro, & Luther, 2014). The benefits of

ω -3FA consumption to prevent mental disorders in pregnant women have not been fully explored neither globally nor among Mexican population. Currently, the study of mental health during pregnancy has been oriented towards psychosocial factors associated to anxiety and depression (Navarrete et al., 2012; Lara et al., 2015), but little is known about the impact of nutrition on these disorders.

A deeper knowledge about the relation of ω -3FA with AS and DS will allow the broadening the range of strategies for preventing and treating these perinatal mental disorders. Hence, the main objective of this study is to evaluate the association between dietary intake of EPA and DHA and anxiety and depressive symptoms in Mexican pregnant women. The study also aims to evaluate the dietary intake of ω -3FA and the prevalence of AS and DS in Mexican pregnant women.

METHOD

Study design and participants

A cross-sectional and analytic study was performed. Between December, 2013 and July, 2014, 687 pregnant women receiving antenatal care were invited to participate. Only 151 of them met the inclusion criteria. The population sample was selected by convenience. Women were approached in the waiting rooms of an urban health center in the city of Morelia, Michoacan, Mexico. This is a community health center that provides prenatal care and other comprehensive medical care to low income women. Invitations to participate were extended to all pregnant women who met inclusion criteria: between 20 and 35 years, in their second trimester of pregnancy and who agreed to participate in the study. We excluded women with multiple pregnancies, acute chronic complications, diagnosed for major depressive disorder, or having consumed supplements with ω -3FA. Finally, women who did not complete the evaluation instruments were eliminated from the sample (Figure 1).

Instruments

The interview included four sections: 1. demographic data; 2. AS; 3. DS; and 4. dietary intake of ω -3FA. A semi-structured interview type survey was performed by undergraduate students in nutrition, who applied all questionnaires; they were previously trained. The research team developed a manual which was given to each of the students to standardize the procedures of the interview.

Demographic data

Age, marital status, education level, occupation, and multiparity.

Anxiety symptoms

AS were assessed by the Trait-Anxiety scale of the Spanish version of the State-Trait Anxiety Inventory (STAI). The STAI consists of the state and trait subscales. We used the trait subscale that measures relatively stable individual propensity to respond with elevated anxiety, consisting of 20 statements in a Likert-type response scale ranging between one and four points. Its internal consistency varies between .85 and .87 (Spielberger & Díaz-Guerrero, 1975). A cut-off > 40 points on both state and trait scales yielded optimal sensitivity (80.95%), specificity (79.75%), positive predictive value (51.05%), and negative predictive value (94%) to determine cases of anxiety in pregnancy (Meades & Ayers, 2011). We used a cut-off > 40 points for determining women with AS, and a > 75 percentile was scored as high AS.

Depressive symptoms

The Edinburgh Postnatal Depression Scale (EPDS), designed by Cox in 1987, was used to evaluate DS suffered during the seven previous days. It was validated in Mexican pregnant women by Alvarado-Esquivel, Sifuentes-Álvarez, Salas-Martínez, and Martínez-García (2006). The EPDS consists of 10 statements with four options about severity/intensity of symptoms rated in a zero to three score. A cut-off score of ≥ 12 was used (Alvarado-Esquivel et al., 2006; Almanza-Muñoz, Salas-Cruz, & Olivares-Morales, 2011). In Mexican women, the EPDS has shown a sensitivity of 75%, a specificity of 93% (Alvarado-Esquivel et al., 2006), and a reliability of $\alpha = .87$ during pregnancy (Rosario-Juárez, Santos-García, Lara-Cantú, & Almanza-Muñoz, 2009).

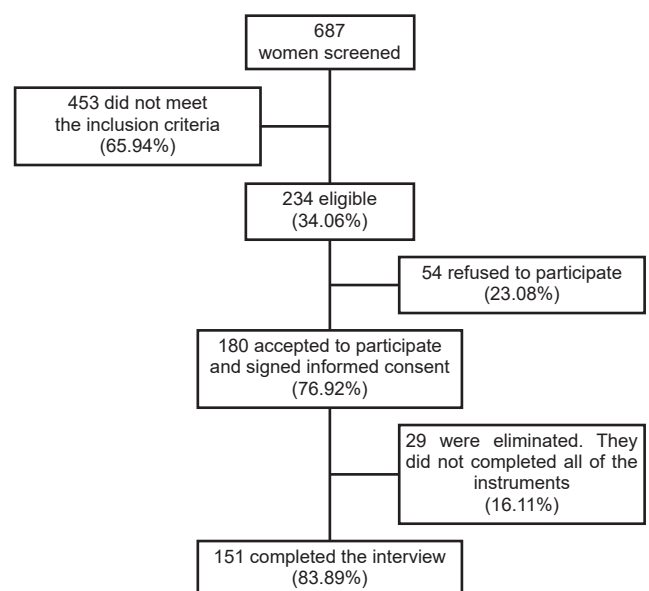


Figure 1. Participants flow from screening to assessment.

Dietary intake of ω -3FA and calculation of nutrient intakes

A semi-quantitative Food Frequency Questionnaire (FFQ) to obtain the average of ω -3FA intake and the nutritional adequacy (NAI) was used. The FFQ was designed by the Instituto Nacional de Salud Pública (INSP) (Hernández-Ávila et al., 1998) and has already been validated in Mexican women residing in Mexico City (Parra et al., 2002). The questionnaire includes 104 items, divided in 10 food sections and an additional section inquiring about the dietary sources of ω -3FA: fresh fish, sardines, tuna, and seafood, and a section of oils. We added extra food items as sources for ω -3FA, such as cod, trout, salmon, and typical dry fish found in México, called *charales*. The questionnaire inquires about the average frequency of consumption of each food item during the previous year. The frequency options of consumption are: never, less than once per month, 1-3 times per month, 1-2 per week, 3-4 per week, 5-6 per week, daily, 2-3 times per day, 4-5 per day, and 6 or more per day. The FFQ has proved to be useful in categorizing women by relative intakes, and can be used as a methodological tool in studies relating maternal diet during pregnancy (Parra et al., 2002). The average daily intake of DHA, EPA, and ALA was calculated using the software System of Evaluation of Nutritional Habits and Consumption of Nutrients (SNUT) v3.0, and developed at the INSP (Hernández-Ávila et al., 1998). SNUT uses the US Department of Agriculture food composition tables to obtain the nutrient content of ω -3FA and its derivatives. Parra et al. (2002) evaluated the validity of this FFQ against a biochemical marker of fat intake, erythrocyte cell membrane phospholipid levels, during pregnancy. They concluded that this FFQ estimates the average long-term intakes of ω -3FA and correlated well with their erythrocyte cell membrane phospholipid status.

Once the average intake of each ω -3FA was calculated, we assessed NAI to observe the degree of dietary insufficiency, comparing the estimated average daily consumption with the adequate intake of each ω -3FA, established by the IOM (Simopoulos, 1999). NAI between 90% and 110% indicates dietary sufficiency, a value between 75% and 89% indicates mild insufficiency, between 74% and 50% indicates moderate insufficiency, and less than 50% stands for severe insufficiency (Córdoba-Caro, Luego, & García, 2012).

Statistical analysis

Using a Kolmogorov-Smirnov test, a non-normal distribution was determined for the daily intake of ω -3FA, and for the anxiety and depression test rates; therefore, non-parametric tests were deemed appropriate. In the descriptive analysis we used frequencies and percentage to determine demographic and psychological characteristics of pregnant women. Categorical variables for establishing insufficient intake of DHA, EPA, and ALA (< 50% of daily recom-

mended intake) were generated. Median, minimum, and maximum values to determine dietary intake and NAI of ω -3FA were used. Spearman rank correlation test and simple linear regression models were generated to evaluate the association between the psychological scales (STAI-T and EPDS) and the dietary intake of DHA and EPA. As a measure of goodness of fit, the coefficient of determination was used. The sample size was calculated with statistic data published (Almanza-Muñoz et al., 2011). A type 1 error of .05 was established, with a confidence interval of 95% ($z = 1.96$). Statistical analysis was conducted in SPSS 20.0.

Ethical considerations

The methodology adhered to the Ethical Principles for Medical Research Involving Human Subjects of the Declaration of Helsinki (World Medical Association, 2013). Informed consent was obtained from each participant. The protocol was evaluated and approved by the Bioethics Committee of the Hospital de la Mujer in Morelia.

RESULTS

Demographic and psychological characteristics are presented in Table 1. The average age of the sample was 25.66

Table 1

Demographic and psychological characteristics of pregnant women ($N = 151$)

	<i>f (%)</i>
Marital status	
Cohabiting	67 (44.4%)
Married	54 (35.8%)
Single	29 (19.2%)
Divorced	1 (.7%)
Education level	
Elementary school	26 (17.22%)
Middle school	54 (35.8%)
High school/equivalent	46 (30.5%)
Bachelor degree/equivalent	25 (16.6%)
Occupation	
Home	93 (61.6%)
Employee	32 (21.2%)
Student	19 (12.6%)
Own business	7 (4.6%)
Multiparity	
Primiparous	52 (34.4%)
Multiparous	99 (65.6%)
Anxiety symptoms (STAI-T) ≥ 40	67 (44.4%)
High anxiety symptoms (STAI-T) > 75th percentile	21 (13.9%)
Depressive symptoms (EPDS) ≥ 12	27 (17.9%)

Note: STAI-T = State-Trait Anxiety Inventory; EPDS = Edinburgh Postnatal Depression Scale.

Table 2
Dietary intake and nutritional adequacy index of omega-3 fatty acid obtained through the Mexican FFQ. (N = 151)

	Dietary intake Median (Min-Max)	Nutritional adequacy index Median (Min-Max)
22:6 ω-3	70 mg/d (0 – 720)	23.30% (0 – 240)
20:5 ω-3	30 mg/d (0 – 280)	13.64% (0 – 127.27)
18:3 ω-3	980 mg/d (0 – 5220)	70.00% (0 – 372.86)
Number and percentage of women with a dietary intake < 50% of daily recommended intake		
22:6 ω-3 ^a	122 (80.8%)	
20:5 ω-3 ^b	138 (91.4%)	
18:3 ω-3 ^c	40 (26.5%)	

Note: FFQ: Food Frequency Questionnaire; ^aDaily recommended intake of DHA: 300 mg/d; ^bDaily recommended intake of EPA: 220 mg/d; ^cDaily recommended intake of ALA: 1400 mg/d.

(SD = 4.50) years. AS were more frequent than DS (44.4% vs. 17.9%, respectively), and 13.9% of the women had high AS (STAI-T > 75th percentile).

Table 2 shows dietary intake and NAI of ω-3FA. Daily intakes of DHA, EPA, and ALA were 70 mg/d, 30 mg/d, and 980 mg/d, respectively; NAI was low for all of the assessed fatty acids. The most insufficient ω-3FA was EPA and its NAI was 13.64%; this indicates that the sample is consuming only 13.64% of daily recommendation of EPA, followed by DHA (23.30%). Likewise, 91.4% and 80.8% of the pregnant women had a severe insufficiency ($\leq 50\%$ of NAI) of EPA and DHA, respectively.

Table 3 shows the Spearman's correlations between DHA and EPA, and scores from STAI-T and EPDS questionnaires. These variables were analyzed for the entire population Spearman's correlations were then performed only for pregnant women who had a severely insufficient intake of DHA and EPA ($n = 122$ and $n = 138$, respective-

Table 3
Correlation between dietary intake of DHA and EPA and STAI-T and EPDS scores

	Spearman's Rho	P value
22:6 ω-3 ^a (N = 151)		
STAI-T scores	-.146	.073
EPDS scores	-.134	.101
20:5 ω-3 ^a (N = 151)		
STAI-T scores	-.175	.032*
EPDS scores	-.142	.082
22:6 ω-3 ^b (N = 122)		
STAI-T scores	-.266	.003*
EPDS scores	-.291	.001**
20:5 ω-3 ^b (N = 138)		
STAI-T scores	-.203	.017*
EPDS scores	-.198	.020*

Note: ^aCorrelation in the entire sample; ^bCorrelation in the group of women who had an insufficient dietary intake (≤ 150 mg DHA/d, ≤ 110 mg EPA/d); * $p < .05$; ** $p < .001$.

ly). In the analysis with the entire sample, a significant negative correlation was found only between intakes of EPA and STAI-T scores ($Rho = -.175$; $p = .032$). For the group of women with severe insufficiency, there was a significant negative correlation between DHA and STAI-T ($p = .003$) and EPDS ($p = .001$) scores. A significant negative correlation with EPA in both mental health scales was also found ($p = .017$ and $p = .020$, respectively).

Figure 2 shows the dispersion models, for these group of women only. To assess the proportion of total variability of the dependent variables (AS and DS), we obtained the coefficient of determination (R^2). In all the dispersion models, R^2 showed a low predictive capacity to the variable, one value on x-axis corresponding to several values in y-axis. The 8.5% and 8.9% of the variability of AS and DS, respectively, can be attributed to a relationship with an intake of DHA; and 2.7% and 1.5%, can be attributed to a relationship with an intake of EPA.

DISCUSSION AND CONCLUSION

In this study, a moderate weak negative correlation between ω-3FA and AS and DS in pregnant women was found, which was significant only when we analyzed women who had an insufficient dietary intake of DHA and EPA. We decided to exclude from correlation's analyses those women who covered or exceeded adequate intake to avoid any bias and reach sound results when analyzing a sample similar to the pregnant Mexican women population. Some cross-sectional studies indicate that people consuming fish and seafood less than once a week have higher risk of developing depressive disorders (Hibbeln, 2002; Markhus et al., 2013; Grosso et al., 2014). However, these results are inconclusive given that it is unknown if one event precedes the other, that is, if eating fish precedes depressive disorders, or if such disorders lead to consuming less fish. In that same context, in cohort studies it has been found that a higher intake of fish together with a higher intake of EPA and DHA were associated with lower rates of DS (Li, Dai, Ekperi, Dehal, & Zhang, 2011; Grosso et al., 2014; Su et al., 2015). Murine models have shown that intake of DHA causes a reduction in the development of behaviors similar to depression (Mizunoya et al., 2013; Su et al., 2015). In humans, Vaz et al. (2013) recorded that a lower intake of ω-3FA and a lower frequency of fish and seafood consumption were associated with higher incidence of AS in pregnant women.

This moderate correlation in our study may imply that, as the insufficiency dietary intake of DHA and EPA in pregnant women becomes more severe, the scores of STAI-T and EPDS increase slightly. It must be stated that in order to observe such a tendency, the intake of EPA and DHA must be severely insufficient ($\leq 50\%$ of the daily recommendation), which seems very likely in Mexico, given that

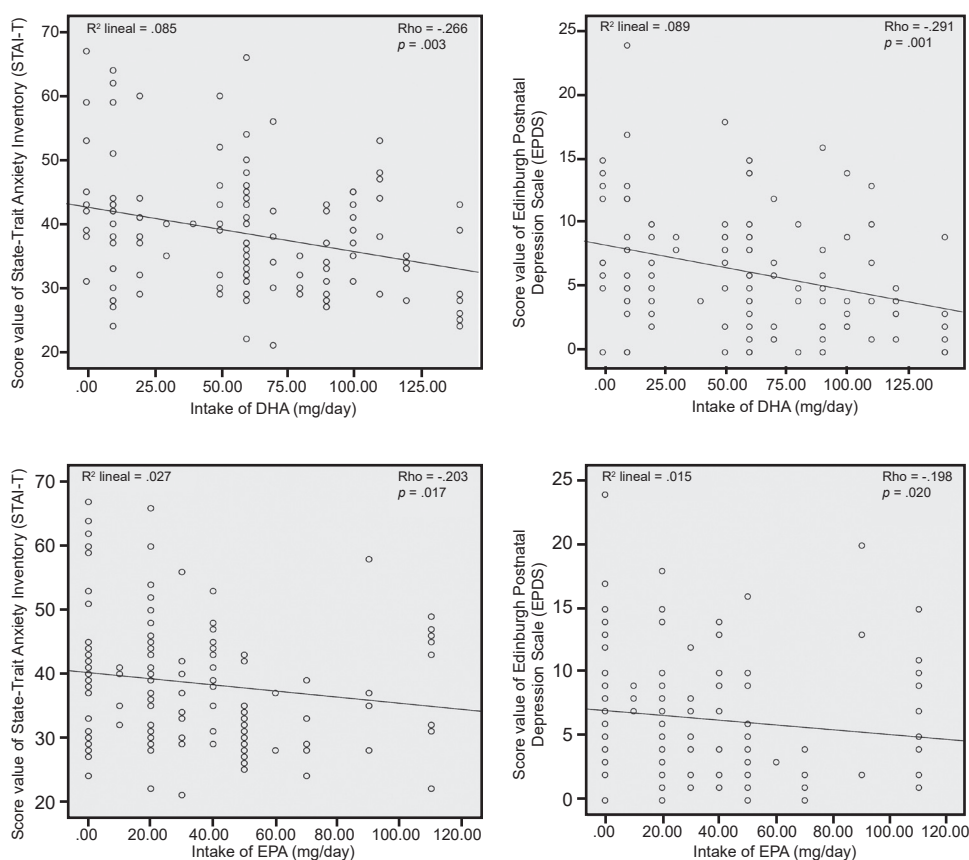


Figure 2. Correlation between intake of ω -3FA (DHA and EPA), and score values of applied scales assessed (STAI-T and EPDS) during the second half of pregnancy in a group of women with insufficient intake.

Mexicans consume fish and seafood approximately once a month (Parra et al., 2002). Nevertheless, considering that the present study is a cross-sectional type, a cause-effect relationship cannot be established. It is suggested to conduct intervention studies in order to assess the efficacy and role played by ω -3FA in AS and DS.

In terms of the studied nutritional variables, it is worrisome that 80.8% and 91.4% of pregnant women had a dietary intake lower than the 50% of the daily recommendation of DHA and EPA, respectively. Such findings coincide with a study from the ENSANUT 2006 that found that the Mexican adult population has an average intake of 300 mg/d of ω -3FA, of which 270 mg correspond to ALA and less than 30 mg correspond to DHA (Ramírez-Silva, Villalpando, Moreno-Saracho, & Bernal-Medina, 2011); in this study, we found intake levels even lower than those. Today, western societies are characterized by a decrease in ω -3FA intake, mainly because EPA and DHA can only be found in fish and seafood products, which are much more expensive than food with other FA, like omega-6 fatty acids. It is worth mentioning that as Morelia is not a coastal city, it would be interesting to compare the dietary intake of ω -3FA, and AS

and DS between non-coastal and coastal regions of Mexico, where people are more likely to consume higher amounts of fish and sea food.

Prevalences of 44.4% and 17.9% in AS and DS, respectively, were found. The 44.4% of women with AS is similar to the prevalence reported in the UK (36% - 56%) and Uzbekistan (38% - 42%) (Meades & Ayers, 2011). However, our results in the prevalence of AS differ considerably from other previous Mexican perinatal studies, which range from 8.1% to 16%, based on self-report questionnaires (Ceballos-Martínez et al., 2010; Navarrete et al., 2012). This figure is in agreement with our results obtained at the cut-off point above the 75th percentile (13.9%). Despite these figures, AS during pregnancy has received little attention (Navarrete et al., 2012). On the other hand, our prevalence of DS was similar to the international report of 18.4% (Gavin et al., 2005), and to the 16.6% prevalence in Mexican pregnant population (Lara et al., 2015). However, our percentage of prevalence is higher than the 12.8% reported by Bennett, Einarson, Taddio, Koren, and Einarson (2004) for the second trimester of pregnancy. These comparisons show that AS and DS are common in the studied population.

There are two limitations in this cross-sectional study. First, we decided to use a convenient non-probabilistic sampling procedure; pregnant women who had enough time and willingness to answer all interview were intentionally selected. Non-randomized sampling limits the scope of generalization to the population as a whole, since some of its characteristics may not be well represented in the sample. The second limitation of the study refers to the waiting rooms, where interviews were conducted, since there was not enough privacy to apply the scales of mood disorders. This situation could lead the participants to responding with a high social desirability for fear of being heard by other people.

These results are promising because they imply that dietary insufficiency of EPA and DHA may be related to the presence of AS and DS. In addition, they reveal that a low intake of DHA and EPA is a predisposing but not triggering factor of AS and DS in pregnant women, suggesting that dietary interventions can be used in order to prevent or reduce the prevalence of AS and DS in pregnant women. It is worth mentioning that women are reluctant to take medication during pregnancy (Lara, 2014), and supplementation with ω -3FA may be a more accepted intervention for AS and DS. Perinatal mental health has been given little attention in Mexico, so these data aim to contribute to increasing evidence on its importance and possible interventions in order to reduce the potential negative effects on women's and infant's physical and mental health. The significant percentage of pregnant women with AS and DS must get the attention of health institutions in order to adopt reliable tools to identify women at risk. This study is the first one in Mexico that evaluates the relationship between mental disorders and nutritional factors such as ω -3FA.

Funding

This study was supported by the Consejo Nacional de Ciencia y Tecnología de México (CONACYT) grant 286473.

Conflict of interests

All authors declare that they have no conflict of interest.

Acknowledgements

The authors wish to thank the undergraduate female students of the Nutrition degree of the UMSNH and UNLA for their technical support during interviews and application of questionnaires to patients.

REFERENCES

Almanza-Muñoz, J., Salas-Cruz, C., & Olivares-Morales, A. (2011). Prevalencia de depresión postparto y factores asociados en pacientes puérperas de la Clínica de especialidades de la Mujer. *Revista de Sanidad Militar*, 65(3), 78-86.

Alvarado-Esquivel, C., Sifuentes-Álvarez, A., Salas-Martínez, C., & Martínez-García, S. (2006). Validation of the Edinburgh Postpartum Depression Scale in a population of puerperal women in Mexico. *Clinical Practice and Epidemiology in Mental Health*, 2(33), 1-5, doi: 10.1186/1745-0179-2-33.

Assies, J., Pouwer, F., Lok, A., Mocking, R., Bockting, C., Visser, I., ... Schene, A. (2010) Plasma and Erythrocyte Fatty Acids Patterns in Patients with Recurrent Depression: A Matched Case-Control Study. *PLOS ONE*, 5(5), 1-9.

Bazan, N., Musto, A., & Knott, E. (2011). Endogenous signaling by omega-3 DHA derived mediators sustains homeostatic synaptic and circuitry integrity. *Molecular Neurobiology*, 44, 216-222.

Bennett, H., Einarson, A., Taddio, A., Koren, G., & Einarson, T. (2004). Depression during pregnancy. Overview of Clinical Factors. *Clinical Drug Investigation*, 24(3), 157-179.

Berenzon, S., Lara, M. A., Robles, R., & Medina-Mora, M. E. (2013). Depresión: estado del conocimiento y la necesidad de políticas públicas y planes de acción en México. *Salud Pública de México*, 55(1), 74-80.

Ceballos-Martínez, I., Sandoval-Jurado, L., Jaimes-Mundo, E., Medina-Peralta, G., Madera-Gamboa, J., & Fernández-Arias, Y. F. (2010). Depresión durante el embarazo Epidemiología en mujeres mexicanas. *Rev Med Inst Mex Seguro Soc*, 48(1), 71-74.

Córdoba-Caro, L. G., Luego, L. M., & García, V. (2012) Adecuación nutricional de la ingesta de los estudiantes de secundaria de Badajoz. *Nutrición Hospitalaria*, 27(4), 1065-1071.

De Castro, F., Place, J. M., Villalobos, A., & Allen-Leigh, B. (2015) Sintomatología depresiva materna en México: prevalencia nacional, atención y perfiles poblacionales de riesgo. *Salud Pública de México*, 57, 144-154.

Ding, Z., Wu, Y., Xu, S., Zhu, R., Jia, X., Zhan, S., ... Tao, F. (2014). Maternal anxiety during pregnancy and adverse birth outcomes: A systematic review and meta-analysis of prospective cohort studies. *Journal of Affective Disorders*, 159, 103-110. doi: 10.1016/j.jad.2014.02.027

Fadzil, A., Balakrishnan, K., Razali, R., Sidi, H., Malapan, T., Japaraj, R. P., ... Manaf, M. R. (2013). Risk factors for depression and anxiety among pregnant women in Hospital Tuanku Bainum, Ipoh, Malaysia. *Asia-Pacific Psychiatry*, 5(1), 7-13. doi: 10.1111/appy.12036.

Field, T., Diego, M., Hernández-Reif, M., Figueiredo, B., Deeds, O., Ascencio, A., ... Kuhn, C. (2010) Comorbid depression and anxiety effects on pregnancy and neonatal outcome. *Infant Behavior & Development*, 33(1), 23-29.

Field, T. (2011) Prenatal depression effects on early development: A review. *Infant Behavior and Development*, 34(1), 1-14. doi: 10.1016/j.infbeh.2010.09.008

Gavin, N. I., Gaynes, B. N., Lohr, K. N., Meltzer-Brody, S., Gartlehner, G., & Swinson, T. (2005) Perinatal depression: a systematic review of prevalence and incidence. *Obstetrics & Gynecology*, 106(5), 1071-1083.

Grosso, G., Galvano, F., Marventano, S., Malaguamera, M., Bucolo, C., Drago, F., & Caraci, F. (2014). Omega-3 fatty acids and depression: Scientific Evidence and Biological Mechanisms. Hindawi Publishing Corporation. *Oxidative Medicine and Cellular Longevity*, 313570, 1-16. doi: 10.1155/2014/313570

Hanley, G., & Oberlander, T. (2014). The effect of perinatal exposures on the infant: antidepressants and depression. *Best Practice & Research Clinical Obstetrics & Gynaecology*, 28(1), 37-48. doi: 10.1016/j.bpobgyn.2013.09.001

Hernández-Ávila, M., Romieu, I., Parra, S., Hernández-Ávila, J. E., Madrigal, H., & Willett, W. (1998). Validity and reproducibility a food frequency questionnaire to assess dietary intake of women living in Mexico City. *Salud Publica de México*, 40(2), 133-140.

Hibbeln, J. R. (2002). Seafood consumption, the DHA content of mother's milk and prevalence rates of postpartum depression: a cross-national, ecological analysis. *Journal of Affective Disorders*, 69(1-3), 15-29.

Hoffmire, C., Block, R., Thevenet-Morrison, K., & van Wijngaarden, E. (2012). Associations between omega-3 poly-unsaturated fatty acids from fish consumption and severity of depressive symptoms. An analysis of the 2005-2008 National Health and Nutrition Examination Survey. *Prostaglandins, Leukotrienes and Essential Fatty Acids*, 86(4-5), 155-160. doi: 10.1016/j.plefa.2012.03.003

Lara, M. A. (2014). Perinatal Depression in Mexican Women: Prevalence, Risk Factors, and Prevention of Postpartum Depression. In: K. Wisner, S. Lara-Cinisomo, (Eds.), *Perinatal Depression Among Spanish-Speaking and Latin American Women. A Global Perspective on Detection and Treatment* (pp. 1-13). New York: Springer. doi: 10.1007/978-1-4614-8045-7

Lara M. A., Navarrete, L., Nieto, L., Barba, J. P., Navarro, J. L., & Lara-Tapia, H. (2015). Prevalence and incidence of perinatal depression and depressive symp-

- toms among Mexican women. *Journal of Affective Disorders*, 175, 18-24. doi: 10.1016/j.jad.2014.12.035
- Li, Y., Dai, Q., Ekperi, L. I., Dehal, A., & Zhang, J. (2011). Fish consumption and severely depressed mood, findings from the first national nutrition follow-up study. *Psychiatry Research*, 190(1), 103-109. doi: 10.1016/j.psychres.2011.05.012
- Liu, J., Green, P., Mann, J., Rapoport, S., & Sublette, E. (2015). Pathways of polyunsaturated fatty acid utilization: Implications for brain function in neuropsychiatric health and disease. *Brain Research*, 1597, 220-246. doi: 10.1016/j.brainres.2014.11.059
- Markhus, M. W., Skotheim, S., Graff, I. E., Frøyland, L., Braarud H. C., Stormark, J. M., & Malde, M. K. (2013). Low Omega-3 Index in Pregnancy Is a Possible Biological Risk Factor for Postpartum Depression. *PLOS ONE*, 8(7), 1-12. doi: 10.1371/journal.pone.0067617
- Meades, R., & Ayers, S. (2011). Anxiety measures validated in perinatal populations: A systematic review. *Journal of Affective Disorders*, 133(1-2), 1-15. doi: 10.1016/j.jad.2010.10.009
- Mizunoya, W., Ohnuki, K., Baba, K., Miyahara, H., Shimizu, N., Tabata, K., ... Ikeuchi, Y. (2013). Effect of dietary fat type on anxiety-like and depression-like behavior in mice. *Springer Plus*, 2(165), 1-9. doi: 10.1186/2193-1801-2-165
- Morse, N. (2012). Benefits of Docosahexaenoic Acid, Folic Acid, Vitamin D and Iodine on Foetal and Infant Brain Development and Function Following Maternal Supplementation during Pregnancy and Lactation. *Nutrients*, 4(7), 799-840. doi: 10.3390/nu4070799
- Navarrete, L. E., Lara, M. A., Navarro, C., Gómez, M. E., & Morales, F. (2012). Factores psicosociales que predicen síntomas de ansiedad postnatal y su relación con los síntomas depresivos en el postparto. *Revista de Investigación Clínica*, 64(6), 625-633.
- Parra, M. S., Schnaas, L., Meydan, M., Perroni, E., Martínez, S., & Romieu, I. (2002). Erythrocyte cell membrane phospholipids levels compared against reported dietary intakes of polyunsaturated FA in pregnant Mexican women. *Public Health Nutrition*, 5(6a), 931-937. doi:10.1079/PHN2002381
- Rahman, A., Fisher, J., Bower, P., Luchters, S., Tran, T., Yasamy, M. T., ... Waheed, W. (2013). Interventions for common perinatal mental disorders in women in low- and middle-income countries: a systematic review and meta-analysis. *Bulletin of the World Health Organization*, 91, 593-601. doi: 10.2471/BLT.12.109819
- Ramírez-Silva, I., Villalpando, S., Moreno-Saracho, J., & Bernal-Medina, D. (2011). Fatty acids intake in the Mexican population. Results of the National Nutrition Survey 2006. *Nutrition & Metabolism*, 8(33), 1-10. doi: 10.1186/1743-7075-8-33
- Rizzo, A. M., Corsetto, P. A., Montorfano, G., Opizzi, A., Faliva, M., Giacosa, A., ... Rondanelli, M. (2012). Comparison between the AA/EPA ratio in depressed and non-depressed elderly females: Omega-3 fatty acids supplementation correlates with improved symptoms but does not change immunological parameters. *Nutrition Journal*, 11(82), 1-11.
- Rosario-Juárez, I., Santos-García, R., Lara-Cantú, M. A., & Almanza-Muñoz, J. J. (2009). Consistencia interna y análisis factorial de la Escala de Depresión Postparto de Edimburgo en mujeres mexicanas embarazadas y puérperas. Reporte preliminar. *Revista Neurología, Neurocirugía y Psiquiatría*, 42(1-4), 1-6.
- Rubertsson, C., Hellström, J., Cross, M., & Sydsjö, G. (2014). Anxiety in early pregnancy: prevalence and contributing factors. *Archives of Women's Mental Health*, 17(3), 221-228. doi: 10.1007/s00737-013-0409-0
- Simopoulos, A. P. (1999). Essential fatty acids in health and chronic disease. *The American Journal of Medical Nutrition*, 70(3), 560s-569s. doi: 10.1093/ajcn/70.3.560s
- Simopoulos, A. P. (2011). Evolutionary Aspects of Diet: The Omega-6/Omega-3 Ratio and the Brain. *Molecular Neurobiology*, 44(2), 203-215. doi: 10.1007/s12035-010-8162-0
- Spielberger, C., & Díaz-Guerrero, R. (1975). *IDARE Inventario de Ansiedad: Rasgo-Estado. Manual e Instructivo*. México: El Manual Moderno, 1-28.
- Su, K. P., Matsuoka, Y., & Pae, C.U. (2015). Omega-3 Polyunsaturated Fatty Acids in Prevention of Mood and Anxiety Disorders. *Clinical Psychopharmacology*, 13(2), 129-137. doi: 10.9758/cpn.2015.13.2.129
- Van Eijsden, M., Hornstra, G., van der Wal, M., Vrijkotte, T., & Bonsel, G. (2008). Maternal ω -3, n-6, and trans fatty acid profile early in pregnancy and term birth weight: a prospective cohort study. *The American Journal of Clinical Nutrition*, 87(4), 887-95.
- Vaz, J., Kac, G., Emmett, P., Davis, J. M., Golding, J., & Hibbeln, J. (2013). Dietary Patterns, ω -3FA Intake from Seafood and High Levels of Anxiety Symptoms during Pregnancy Findings from the Avon Longitudinal Study of Parents and Children. *PLOS ONE*, 8(7), 1-9. doi: 10.1371/journal.pone.0067671
- Wisner, K. L., Lara-Cinisomo, S., Pinheiro, E., & Luther, J. (2014). Characteristics of Hispanic Women Screened for Postpartum Depression. In: K. Wisner, S. Lara-Cinisomo (Eds.) *Perinatal Depression Among Spanish-Speaking and Latin American Women. A Global Perspective on Detection and Treatment* (pp. 1-13), New York: Springer. doi: 10.1007/978-1-4614-8045-7_1
- World Medical Association. (2013). Declaración de Helsinki. Principios éticos para las investigaciones con seres humanos. 64ª Asamblea General, Fortaleza, Brasil, octubre de 2013. Retrieved from: <https://www.wma.net/es/policies-post/declaracion-de-helsinki-de-la-amm-principios-eticos-para-las-investigaciones-medicas-en-seres-humanos/>