Vitamin D Deficiency and Depression in Obese Adults: A Case-Control Study

Leila Kamalzadeh
Iran University of Medical Sciences

Atefeh Ghanbari Jolfaei (ghanbari.a@iums.ac.ir)
Iran University of Medical Sciences

Malihe Saghaei
Iran University of Medical Sciences

Research Article

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Abstract

Background: Depression is one of the most prevalent psychiatric disorders reported in obese population. Amongst the contributing factors of depression, vitamin D deficiency has increasingly drawn attention in recent years. This paper seeks to examine the association between serum vitamin D level and depression in patients with obesity.

Methods: This case-control study included 173 depressed obese patients and 174 non-depressed controls. Structured Clinical Interview for DSM-5 (SCID-5) was used to confirm the diagnosis of depression. 25-Hydroxyvitamin D [25(OH)D], Thyroid stimulating hormone (TSH), fasting blood sugar (FBS), parathyroid hormone (PTH) levels, and BMI were assessed in both groups. The statistical analyses included T-test, Chi-squared test, and multivariable logistic regression.

Results: The mean 25(OH)D levels were significantly different between the case and control groups (20.43 ± 15.37 vs. 26.55 ± 13.17, P < .001). Vitamin D insufficiency/deficiency was detected in 77.6% and 67.4% of the case and control groups, respectively, which was significantly different (P = .034). Being female, greater age and lower vitamin D levels were associated with greater odds of developing depression (OR = 3.57, 95% CI = 1.82-7.02; OR = 1.05, 95% CI = 1.02-1.07; OR = 1.51, 95%CI = 1.16-1.96, respectively).

Conclusion: The present study provides additional evidence with respect to the hypothesis that low vitamin D serum concentration is associated with depression in obese adults, and highlights the need for further research to determine whether this association is causal.

Introduction

Obesity is increasingly recognized as a serious public health concern. More than 650 million people throughout the world suffer from overweight and obesity(1). There is strong evidence to suggest that obesity is conjoint with several mental and physical problems(2). Depression is one of the most prevalent psychiatric disorders reported in obese population contributing to significant disability, mortality and healthcare costs(3). The exact mechanisms linking depression and obesity have not been established. Studies have shown multiplex interactions between biologic, psychologic, and environmental factors giving rise to the association between obesity and depression(4). Amongst the contributing factors of depression, vitamin D deficiency has increasingly drawn attention in recent years(5).

Vitamin D, also known as cholecalciferol, is a unique neuro-steroid hormone which is vital for numerous brain functions. This hormone binds to receptors in numerous regions of the brain including the hippocampus and cingulate cortex, which are involved in the pathogenesis of depression and other mental illnesses(6). Many clinical studies have found depression, anxiety and cognitive impairment to be associated with low serum levels of 25-hydroxyvitamin D [25(OH)D], which is the major circulating form of vitamin D, in average weight people(5–7). Recent research has also provided evidence for antidepressant properties of vitamin D supplementation(8, 9).
Strong evidence of aberrations in the vitamin D-endocrine system as well as low serum 25(OH)D levels have been seen in obese individuals\(^\text{10, 11}\). It has been demonstrated that vitamin D deficiency is 35\% more likely in obese people compared to the healthy-weight subjects\(^\text{12}\). Volumetric dilution of Vitamin D is the most plausible mechanism of low 25(OH)D in individuals with obesity\(^\text{11}\). Alternative mechanisms for lower 25(OH)D concentrations in patients with obesity include lower dietary consumption, decreased dermal synthesis, reduced intestinal absorption, altered metabolism, as well as less sunlight exposure due to lower physical activity\(^\text{13}\).

In spite of a biologically potential role of vitamin D in the development of depression, very few studies have investigated this association in overweight and obese subjects, with conflicting findings. While two studies indicated the benefits of vitamin D supplementation on depressive symptoms in obese adults\(^\text{14, 15}\), a recent randomized trial presented contradictory results\(^\text{16}\).

The discrepancy among these results has given rise to the need for further research. In this regard, this study seeks to examine the association between serum vitamin D level and depression in obese patients.

**Methods**

**Study design**

This case-control study was conducted at Rasoul-e Akram hospital, an affiliate of Iran University of Medical Sciences located in Tehran, Iran.

**Study Participants**

The study population consisted of males and females aged 18 to 60 years old visiting out-patient obesity clinic from April 2019 to October 2020. 173 depressed patients (cases) and 174 non-depressed controls were enrolled in the study. Eligibility criteria required patients to have a BMI greater than or equal to 30, normal thyroid function, normal fasting blood sugar, normal PTH level, no history of sleep apnea, neither past nor present substance use, no history of calcium supplement or vitamin D use. Additional criteria for the non-depressed group included: not meeting criteria for another mental disorder and no history of psychiatric medication use. Patients with incomplete medical records were excluded from the study.

**Measurements**

**Anthropometric Parameters**

Height and weight were measured and body mass index (BMI) was calculated as weight/height\(^2\) (kg/m\(^2\)).

**Assessment of depression**

To confirm the diagnosis of depressive disorder, participants were interviewed by expert psychiatrists using the Structured Clinical Interview for the Diagnostic and Statistical Manual of Mental Disorders (DSM-5) (SCID-5)\(^\text{17, 18}\).
Demographic and clinical information

Participants’ medical information was collected using hospital database which included data on basic demographics, medical and psychiatric diagnoses, history of substance and medication use, and physical and psychiatric examination, as well as laboratory data including serum levels of thyroid stimulating hormone (TSH), fasting blood sugar (FBS), and parathyroid hormone (PTH). Considered laboratory reference ranges were: TSH (0.35–5.5 mIU/L), FBS (< 100 mg/dl), and PTH (0–55 pg/mL).

Vitamin D Assays

The circulating levels of 25(OH)D in the plasma samples were measured using Enzyme immunoassay (EIA) method (IDS, UK). The final measurements were classified according to definitions established by the Endocrine Society: 30–100 ng/ml was considered normal, with insufficient 25(OH)D levels or hypovitaminosis D sub grouped into two: vitamin D insufficiency (20–29 ng/ml) and vitamin D deficiency (< 20 ng/ml)(19).

Statistical analysis

The Statistical Package for Social Sciences, version 22 was used for all statistical analyses (SPSS Inc., Chicago, IL, USA). Statistical significance was analyzed using analysis of variance, T-test and Chi-squared test as appropriate. Multivariable logistic regression was performed to assess the impact of confounders. A $p$-value of < 0.05 was considered statistically significant.

Results

A total of 174 depressed patients and 173 non-depressed controls were studied. Statistically significant differences were observed among the two groups in terms of the ratio of females to males ($P < .001$) and age ($P < .001$), i.e., the case group had a significantly higher proportion of females (92% vs. 75.1%) and a higher mean age than the control group (41.76 ± 10.016 vs. 37.05 ± 9.857). The mean vitamin D levels were also significantly different between the case and control groups (20.43 ± 15.37 vs. 26.55 ± 13.17, $P < .001$). Moreover, vitamin D insufficiency/deficiency was detected in 77.6% and 67.4% of the case and control groups, respectively, which was significantly different ($P = .034$). The mean BMI was not significantly different between the groups ($P = .051$). The baseline characteristics of the participants are summarized in Table 1.
Table 1: Baseline characteristics of the participants

<table>
<thead>
<tr>
<th>Characteristic</th>
<th>Cases</th>
<th>Controls</th>
<th>P value</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>N = 174</td>
<td>N = 173</td>
<td></td>
</tr>
<tr>
<td>Age (years)</td>
<td>41.76 ± 10.016</td>
<td>37.05 ± 9.857</td>
<td>&lt; .001</td>
</tr>
<tr>
<td>Female, n (%)</td>
<td>160 (92.0%)</td>
<td>130 (75.1%)</td>
<td>&lt; .001</td>
</tr>
<tr>
<td>BMI (Kg/m$^2$)</td>
<td>41.93 ± 7.42</td>
<td>43.45 ± 7.02</td>
<td>.051</td>
</tr>
<tr>
<td>Vitamin D level (ng/mL)</td>
<td>20.43 ± 15.37</td>
<td>26.55 ± 13.17</td>
<td>&lt; .001</td>
</tr>
<tr>
<td>Vitamin D status, n (%)</td>
<td>39 (22.4%)</td>
<td>56 (32.3%)</td>
<td>.034</td>
</tr>
<tr>
<td>Normal</td>
<td>135 (77.6%)</td>
<td>117 (67.4%)</td>
<td></td>
</tr>
<tr>
<td>Insufficient/Deficient</td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

Table 2 shows the logistic regression results in which, depression is the dependent variable and age, gender and 25(OH)D are independent variables. As can be seen, vitamin D level, age, and gender are significantly associated with depression.

<table>
<thead>
<tr>
<th>Variable</th>
<th>B</th>
<th>Standard Error</th>
<th>Wald Statistic</th>
<th>P value</th>
<th>OR, 95% CI</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age</td>
<td>.049</td>
<td>.012</td>
<td>16.985</td>
<td>&lt; 0.001</td>
<td>1.05 (1.02–1.07)</td>
</tr>
<tr>
<td>Gender (Being female)</td>
<td>1.275</td>
<td>.344</td>
<td>13.702</td>
<td>&lt; 0.001</td>
<td>3.57 (1.82–7.02)</td>
</tr>
<tr>
<td>Vitamin D level (Insufficient/deficient)</td>
<td>.415</td>
<td>.133</td>
<td>9.783</td>
<td>.002</td>
<td>1.51 (1.16–1.96)</td>
</tr>
<tr>
<td>Constant</td>
<td>-5.308</td>
<td>.916</td>
<td>33.591</td>
<td>&lt; 0.001</td>
<td>.005</td>
</tr>
</tbody>
</table>

Table 2. Multivariable logistic regression analysis including confounders of age and gender

The estimate for the effect of age was .049 (SE = .012), P < 0.001. This translates into an OR of 1.05 (95% CI = 1.02–1.07). This means that an increase in age by 1 year increases the odds of developing depression by 5%.

The estimate for the effect of being female was 1.275 (SE = .344), P < 0.001. This translates into an OR of 3.57 (95% CI = 1.82–7.02). This means that obese females are at 3.57 times greater odds of developing depression.

The estimate for the effect of vitamin D level was .415 (SE = .133), P = .002. This translates to an OR of 1.51 (95% CI = 1.16–1.96). Thus, for each unit decrease in vitamin D level, the odds of developing
Depression increase by a factor of 1.51.

Discussion

The findings of the present study revealed that there is an association between low 25(OH)D serum levels and incident depression in obese patients. The relation between low levels of vitamin D and depression was already established in normal weight individuals by past research (5–7). However, relatively few studies have evaluated this relationship in obese adults, and results have been mixed. Consistent with our findings Jorde et al. (15) and Milaneschi et al. (4) found that low serum level of vitamin D is a risk factor for depression and suggested that BMI plays an important mediating role in the association between vitamin D and depression. In the same vein, Irandoust et al. (14) demonstrated that both vitamin D supplementation and physical activity have beneficial impact on depressive symptoms in obese females. Penckofer et al. (20) yielded comparable results in their study on women with type 2 diabetes. They demonstrated that vitamin D supplementation significantly ameliorated depression in the patients.

Multiple mechanisms are involved in the interaction between vitamin D deficiency, obesity and depression. It has been proposed that an inverse relationship exists between vitamin D serum levels and BMI (21). As was pointed out in the introduction to this paper, low serum 25(OH)D in obese patients can occur for a number of reasons including insufficient vitamin D consumption, increased fat or muscle mass, genotype variation in vitamin D binding proteins or enzymes responsible for vitamin D metabolism (11, 13). On the other hand, vitamin D deficiency can increase the risk of developing depression, through several biological pathways including effects on immunomodulation, cellular signaling, modulation of hypothalamic-pituitary-adrenal axis, intracellular calcium homeostasis, and production of neurotransmitters (6, 22). Moreover, both obesity and vitamin D deficiency lead to a chronic low-grade inflammation, which has been suggested to contribute to the development of depression (11, 23). At last, it is also possible that PTH levels contribute to the relation between vitamin D, BMI, and depression. Recent evidence suggests that both obesity and low vitamin D levels are accompanied by significantly higher PTH levels, and high PTH levels are related to depression (4).

However, more recently, literature has emerged that offers contradictory findings about these associations. In contrast to the mentioned studies, in a randomized controlled study, Mousa et al. (16) showed that depressive symptoms in obese individuals were not associated with 25(OH)D concentrations, nor did improve by Vit D supplementation. Similarly, a recent clinical interventional cohort study revealed that vitamin D supplementation for six months had no significant impact on depressive symptoms, but could improve anxiety symptoms in depressed patients with vitamin D deficiency. The authors suggested that BMI is an important mediating factor between low serum 25(OH)D and anxiety symptoms (24).

Different findings may be attributed to the differences in the study population (different races, gender, and age groups), diverse methodology (different vitamin D supplementation doses and duration) and different baseline levels of serum 25(OH)D concentrations. Moreover, the mentioned studies applied self-
reported psychiatric rating scales for evaluation of depression rather than a clinician-rated assessment, and therefore were prone to multiple potential biases. In addition, some of the previous studies did not adjust for potential confounders, such as thyroid dysfunction, comorbid diabetes mellitus, sleep apnea, low physical activity, and substance use that may play a role in developing depression in obese population(25).

The current study also found that obese females are at greater odds of developing depression compared to obese males. Moreover, the analyses revealed a statistically significant relationship between increased age and developing depression. These results match those observed in earlier studies(26, 27). Several factors may be responsible for this gender difference, including biologic aspects such as genetic features, endocrine system, functions of neurotransmitters, as well as psychosocial variables such as lower appearance satisfaction and Gender-role attitudes(26).

Conclusions

In summary, the present study provides additional evidence with respect to the hypothesis that low vitamin D serum concentration is associated with depression in obese adults, and highlights the need for further research to find out whether this association is causal.

Although the present study is based on a small sample size, the results indicate that low serum vitamin D level can be a modifiable risk factor for depression in obese population. The key strength of this study is the exclusion of several potential confounders such thyroid dysfunction, comorbid diabetes, hyperparathyroidism, and substance use which were not addressed by previous studies. Moreover, in the present study, the diagnosis of depression was based on standardized semi-structured interviews by expert psychiatrists rather than self-report rating scales.

Abbreviations

BMI: Body Mass Index

DSM-5: Diagnostic and Statistical Manual of Mental Disorders

EIA: Enzyme Immunoassay

FBS: Fasting Blood Sugar

25(OH)D: 25-Hydroxyvitamin D

OR: Odds Ratio

PTH: Parathyroid Hormone

SCID-5: Structured Clinical Interview for DSM-5
Declarations

*Ethics approval and consent to participate*

The principles of the World Medical Association Declaration of Helsinki were adopted in the present study (28). This research was approved by the independent ethics committee of Iran University of Medical Sciences (IR.IUMS.REC 1395.8721215026). All patients signed informed consent statements.

*Consent for publication*

Written informed consent was obtained from the patients for publication of this research and any accompanying images. A copy of the written consent is available for review by the Editor-in-Chief of this journal.

*Availability of data and materials*

The datasets used and/or analyzed during the current study are available from the corresponding author on reasonable request.

*Competing interests*

The authors declare that they have no competing interests.

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*Authors’ contributions*

LK have drafted the work and substantively revised it. AGhJ have made substantial contributions to the conception of the work. MS was a major contributor in writing the manuscript. All authors read and approved the final manuscript.

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