

# Assessment of vitamin D levels in correlation with coronary CT angiography results in a Slovak Population

---

## Keywords

vitamin D, angiography, computed tomography, coronary artery disease

---

## Abstract

### Introduction

The growing interest in Vitamin D deficiency as a potential contributor to coronary artery disease (CAD) has prompted increased scrutiny. Nevertheless, its status as a confirmed risk factor remains unestablished. This study seeks to explore the connection between serum vitamin D levels and CAD.

### Material and methods

Employing a cross-sectional approach, 205 patients eligible for coronary computed tomography angiography (CCTA) were chosen. Serum vitamin D levels were assessed and juxtaposed with the outcomes of CCTA, which encompassed the coronary artery calcium score (CACS), as well as the presence and severity of coronary artery involvement attributed to atherosclerotic plaques.

### Results

The average age of the participants was  $61.4 \pm 10.8$  years, and the mean serum vitamin D level was  $19.6 \pm 10.3$  ng/dl (ranging from 4.7 to 39.7 ng/dl). In total, 52.1% of the participants (n=107) exhibited vitamin D deficiency, 35.1% (n=72) had insufficient levels, and 12.6% (n=26) had sufficient levels of vitamin D. The mean serum vitamin D level was notably lower in patients with severe coronary artery disease (CAD) ( $P < 0.0001$ ). According to the Spearman test, a significant negative correlation (-0.48) was identified between the serum vitamin D level and CACS ( $P < 0.0001$ ). Conversely, the mean CACS in the vitamin D deficient group was significantly higher than in the insufficient and sufficient vitamin D groups ( $P < 0.0001$  for both comparisons).

### Conclusions

There was a correlation between vitamin D deficiency and both CACS and the severity of coronary artery stenosis.

## **Assessment of vitamin D levels in correlation with coronary CT angiography results in a Slovak Population**

Dominik Pella<sup>1</sup>, Tibor Poruban<sup>1</sup>, Jozef Gonsorcik<sup>1</sup>, Jan Fedacko<sup>3</sup>, Zuzana Pella<sup>2</sup>, Karolina Angela Sieradzka Uchnar<sup>1</sup>, Radka Hreskova<sup>1</sup>, Marianna Barbierik Vachalcova<sup>1</sup>

<sup>1</sup>East Slovak Institute of Cardiovascular Diseases and School of Medicine, Pavol Jozef Safarik University, Kosice, Slovakia

<sup>2</sup>Department of Cybernetics and Artificial Intelligence, Faculty of Electrical Engineering and Informatics, Technical University of Kosice, Kosice, Slovakia

<sup>3</sup>Gerontology and Geriatrics Clinic, Pavol Jozef Safarik University, Kosice, Slovakia

### **Corresponding author:**

Tibor Poruban, East Slovak Institute of Cardiovascular Diseases and School of Medicine, Pavol Jozef Safarik University, Kosice, Slovakia

Email: tporuban@vusch.sk

Telephone: +421 55 789 1450

### **DECLARATIONS**

Ethics approval and consent to participate - All procedures followed were in accordance with the ethical standards of the responsible committee on human experimentation (IRB/ERC VUSCH, Reference Number 27052021) and with the Helsinki Declaration of 1975, as revised in 2008

Consent for publication - Not applicable

Availability of data and materials - Not applicable

Competing interests - The authors declare that they have no competing interests

Funding - Not applicable

Contributions - All authors contributed to the design and implementation of the research, to the analysis of the results and to the writing of the manuscript

Clinical trial registration - Not applicable

Acknowledgements - The authors would like to acknowledge the support staff for its timely effort, sincerity, and dedication towards work

## **Abstract**

**Introduction:** The growing interest in Vitamin D deficiency as a potential contributor to coronary artery disease (CAD) has prompted increased scrutiny. Nevertheless, its status as a confirmed risk factor remains unestablished. This study seeks to explore the connection between serum vitamin D levels and CAD.

**Material and methods:** Employing a cross-sectional approach, 205 patients eligible for coronary computed tomography angiography (CCTA) were chosen. Serum vitamin D levels were assessed and juxtaposed with the outcomes of CCTA, which encompassed the coronary artery calcium score (CACS), as well as the presence and severity of coronary artery involvement attributed to atherosclerotic plaques.

**Results:** The average age of the participants was  $61.4 \pm 10.8$  years, and the mean serum vitamin D level was  $19.6 \pm 10.3$  ng/dl (ranging from 4.7 to 39.7 ng/dl). In total, 52.1% of the participants (n=107) exhibited vitamin D deficiency, 47.9% (n=98) had no deficient levels of vitamin D. The mean serum vitamin D level was notably lower in patients with severe coronary artery disease (CAD) ( $P < 0.0001$ ). According to the Spearman test, a significant negative correlation (-0.48) was identified between the serum vitamin D level and CACS ( $P < 0.0001$ ). Conversely, the mean CACS in the vitamin D deficient group was significantly higher than in the insufficient and non-insufficient vitamin D groups ( $P < 0.0001$  for both comparisons).

**Conclusions:** There was a correlation between vitamin D deficiency and both CACS and the severity of coronary artery stenosis.

**Keywords:** coronary artery disease, vitamin D, computed tomography, angiography

## Cover Letter

May 12, 2024

Maciej Banach, *Editor-in-Chief*  
Collective of Editorial Office

Dear Editor-in-Chief,

thank you for your letter on our manuscript No. AMS-16606-2024-01 entitled "*Assessment of vitamin D levels in correlation with coronary CT angiography results in a Slovak Population*".

We were pleased to see that the interest rating was encouraging and following your letter we revised our paper to address the helpful and constructive comments. We herewith submit the revised version hoping that the concerns have been adequately addressed and look forward to receiving your response.

Sincerely,

Tibor Poruban, MD, PhD, MPH, MBA

East Slovak Institute of Cardiovascular Diseases and School of Medicine, Pavol Jozef Safarik University,

Ondavska 8, Kosice, Slovakia

E-Mail: poruban.tibor@gmail.com

Telephone: +421 55 789 1450

Preprint

**Response to the editor:**

*1. The authors have improved the quality of the work on several respects. I do recommend still that the authors show at test of the hypothesis testing the hypothesis: “is there an independent association between Vitamin D levels and coronary calcium scores, when adjusted for traditional cardiovascular risk factors?”. This will make the study more robust. Thank you for the opportunity to review this interesting study.*

Dear Editor, we referred to the comment of the reviewer and specified, in more detail. In the Results section, page 11, paragraph Vitamin D, the last sentence was edited as follows: Patients without hyperlipidemia and with lower CACS exhibited significantly higher levels of vitamin D compared to those with hyperlipidemia and with higher CACS ( $p < 0.01$ ) (Table 3). Then, page 12, Table 3, an independent association between Vitamin D levels and coronary calcium scores was tested and confirmed using multivariable regression analysis as shown in a last line of the table.

Preprint

## Introduction

While vitamin D has conventionally been linked to bone health due to its vital involvement in bone and mineral metabolism, it is now being recognized for various non-skeletal functions.<sup>1</sup> Recently, a more extensive role has been suggested for this essential vitamin, attributed to the presence of its receptor in diverse tissues, including cardiovascular organs. Atherosclerosis, a prominent feature of cardiovascular disorders, is a complex process that can evolve over decades before becoming clinically evident.<sup>2</sup> Inflammatory alterations, osteogenesis, angiogenesis, and endothelial dysfunction are among the crucial factors associated with the pathogenesis of atherosclerosis, all of which are either directly or indirectly influenced by vitamin D.<sup>3</sup>

Numerous studies have investigated the correlation between vitamin D status and various aspects of cardiovascular disorders, such as metabolic syndrome, hypertension, or fatal events associated with coronary artery disease (CAD).<sup>4,5</sup> In light of these findings, there is a growing call for heightened attention to the potential role of vitamin D deficiency as a risk factor for CAD. Nevertheless, the existing evidence remains inconclusive due to conflicting reports in the literature, with some supporting and others opposing this concept.<sup>6,7</sup> Importantly, the genetic polymorphism of the vitamin D receptor may alter its role, and variations in ethnicity, lifestyle, and geographic conditions could potentially influence its effects.<sup>8</sup>

The objective of this study was to assess the correlation between vitamin D levels and the findings of coronary artery computed tomography (CT) angiography in a Slovakian population.

## Material and Methods

In this cross-sectional study, we extended invitations to patients referred for coronary artery CT angiography (CTA) by a cardiologist. Exclusions encompassed individuals with recent coronary heart disease, such as prior myocardial infarction, coronary stent or bypass grafting, or recent invasive coronary angiography confirming coronary vessel stenosis. Additionally, those with vitamin D intake, chronic diseases (including known hepatic, renal, endocrine, or metabolic conditions), a documented presence of supraventricular tachyarrhythmia with numerous extrasystoles in resting electrocardiogram (ECG), and individuals allergic to contrast agents were excluded. Informed consent was obtained from all participants following a thorough explanation and clarification of the study procedures.

The sample comprised 205 patients (20 were excluded). Initially, the age and sex of each patient was recorded, as well as any family history of CAD and cigarette smoking. Patients were introduced to a constant laboratory environment at our hospital on the same day of CTA. Blood samples were taken and analyzed, and serum levels of 25-hydroxyvitamin D (25(OH)D) were measured using high-performance liquid chromatography. Based on the levels of vitamin D measured, patients were categorized into two groups as follows: vitamin D deficiency (<20 ng/ml), and non-insufficient level of vitamin D (>20 ng/ml).<sup>9</sup>

All participants underwent a two-phase imaging protocol, beginning with a noncontrast scan followed by contrast-enhanced coronary CT angiography (CTA), utilizing a 64-slice scanner system (SOMATOM Force; Siemens AG, Berlin, Germany). The Agatston scoring method (Aquarius iNtuition software, TeraRecon, Inc., San Mateo, CA, USA) was employed to determine the coronary artery calcium score (CACS), which was recorded for each patient. Patients were then categorized into five groups based on their scores according to Rumberger's guidelines: normal (CACS = 0), minimal (CACS = 1–10), mild (CACS = 11–100), moderate (CACS = 101–400), and severe (CACS > 400).<sup>10</sup> A retrospective electrocardiogram-gated technique was employed, and the scan delay was calculated using the test bolus technique. Intravenously, a bolus of 70 ml of iodinated contrast (Visipaque, GE Healthcare)

was injected, followed by saline, at a flow rate of 4.5 ml/s. The scan parameters included collimation of  $64 \times 0.625$  mm, tube current of 400–600 mA, tube voltage of 120 kV, and gantry rotation time of 350 ms.

Initially, source images were reconstructed at 75% of the R-R interval to serve as default images; however, alternative phase locations were also available and reviewed as needed. Thin slice maximum intensity projections and curved multiplanar reconstructions were employed. The evaluation included the assessment of coronary artery stenosis, with the severity estimated as the ratio of the luminal diameter in the most stenotic area to that of the unaffected distal portion. A stenosis was deemed significant if it reached 50% or more. An independent review of all scans was conducted by both a cardiologist and a radiologist with expertise in coronary CTA. Subsequently, patients were classified into two groups: normal and nonsignificant disease (absence of plaque or presence of plaque with less than 50% stenosis), and significant disease (plaque with more than 50% stenosis in one, two, or three major coronary arteries, respectively).

Quantitative variables, such as continuous measurements, and qualitative variables, representing categorical data, were meticulously detailed by presenting their mean values along with their respective standard deviations. Additionally, proportions were meticulously expressed as percentages to provide a comprehensive understanding of the data distribution. For statistical inference, a robust suite of tests and analyses was employed, including the Kolmogorov-Smirnov test and the Shapiro-Wilk test, to assess the normality of the data distribution. The Spearman correlation coefficient was utilized to explore potential relationships between variables, particularly examining associations between serum vitamin D levels and other pertinent factors. Furthermore, the study conducted multivariable regression analysis, which incorporated a comprehensive set of covariates such as age, gender, smoking status, arterial hypertension, and/or hyperlipidemia. This analytical approach allowed for a thorough investigation into the collective impact of these factors on serum vitamin D levels. To ascertain statistical significance, a stringent two-sided  $\alpha$  level of 0.05 was meticulously applied, ensuring rigorous scrutiny of the findings. The analysis was meticulously



executed using the R Statistics software, version 3.5, an open-source project renowned for its robustness and versatility, which facilitated comprehensive data exploration and interpretation across all participant data sets.

Preprint

## Results

### Clinical characteristics of patients

Out of 225 patients initially considered for the study, 205 met the inclusion criteria. The average age of participants was  $61.4 \pm 10.8$  years, ranging from 30 to 85 years. The age distribution was as follows: 14.1% (n=29) were under 50 years old, 28.3% (n=58) were between 50 and 59 years, 36.1% (n=74) were between 60 and 69 years, and 21.5% (n=44) were 70 years or older. Approximately 43% (n=88) were male, and 16.6% (n=34) were smokers. In terms of the presence of other selected risk factors (arterial hypertension, type 2 diabetes mellitus, elevated LDL-C), their prevalence was as follows. Out of the total of 205 patients, 168 patients were treated for hypertension (80.5%), and 18 patients (8.8%) had documented type 2 diabetes mellitus either on diet, oral antidiabetic therapy, or insulin therapy. The average level of LDL-C was 3.1 mmol/l, with the highest at 5.6 mmol/l and the lowest at 1.34 mmol/l. The average serum vitamin D level was  $19.6 \pm 10.3$  ng/dl, ranging from 4.7 to 39.7 ng/dl. According to the results, 52.1% of all participants (n=107) had, and 47.2% (n=98) had no vitamin D deficiency and were divided into these two groups (**Table 1**).

**Table 1** Summarized characteristics of participants

Determinant	Vit D def. (n = 107)	No Vit D def. (n = 98)	P-value
Age (years)	62.6±11.1	60.9±10.5	0.469
Gender (male, n)	42 (39.3%)	46 (46.9%)	0.520
BMI (kg/cm <sup>2</sup> )	25.8±4.6	27.2±5.7	0.092
Family history with CAD	59 (55.1%)	51 (52%)	0.423
Smokers (n)	26 (24.3%)	12 (12.2%)	0.317
Arterial hypertension (n)	90 (84.1%)	78 (79.5%)	0.069
Type 2 diabetes mellitus (n)	7 (6.5%)	11 (11.2%)	0.654
LDL cholesterol (mmol/l)	2.71±0.31	2.39±0.28	0.030
HDL cholesterol (mmol/l)	0.89±0.18	0.91±0.16	0.663
Total cholesterol (mmol/l)	3.6±0.4	3.4±0.4	0.280
Atherosclerotic CVD risk	12.8±1.8	12.2±1.6	0.826

*BMI, body mass index; CAD, coronary artery disease; HDL, high density lipoprotein; LDL, low density lipoprotein*

### Coronary artery disease

The CT angiographic findings among participants were as follows: 77% (n=158) had normal or nonsignificant CTA; 23% (n=45) had at least one significant coronary artery stenosis.

### Calcium score

Among the participants, 37.5% (n=77), 10.2% (n=21), 17.5% (n=36), 18.5% (n=38), and 17.5% (n=36) had normal, minimal, mild, moderate, and severe calcium scores, respectively. As anticipated, the mean calcium score was notably higher in patients with severe CAD (P<0.0001). The Spearman test revealed a significant negative correlation coefficient (-0.48) between serum vitamin D levels and (CACs) (P<0.0001) (**Figure 2**).

Conversely, the mean CACS in the vitamin D deficient group was significantly higher than in the vitamin D insufficient and non-insufficient groups ( $P<0.0001$ ).

### **Vitamin D**

There was a significant decline ( $P<0.0001$ ) in the mean serum vitamin D level with increasing severity of CAD (**Figure 1**). In the logistic regression analysis, among the determinants including age, gender, smoking habits, arterial hypertension, hyperlipidemia, and serum vitamin D level, the latter emerged as a significant determinant of CAD (**Table 2**). This implies that individuals with adequate vitamin D levels have a lower cardiovascular risk and are less likely to exhibit hemodynamically significant stenosis. Conversely, those with a vitamin D deficiency fall into a higher cardiovascular risk category, and a greater likelihood of coronary artery disease has been observed in this group. Finally, among the determinants mentioned above, only hyperlipidemia was a significant factor affecting serum vitamin D levels. Patients without hyperlipidemia and with lower CACS exhibited significantly higher levels of vitamin D compared to those with hyperlipidemia and with higher CACS ( $p<0.01$ ) (**Table 3**).

**Table 2** Results of the logistic regression analysis considering age, gender, smoking habits, arterial hypertension, hyperlipidemia, and serum vitamin D level as determinants of CAD

Determinant	OR	95% CI	P-value
Age (years)	0.005	-0.023 - 0.029	0.89
Gender (M/F)	-0.410	-0.814 - 0.123	0.18
Smoking (No/Yes/Ex-smokers)	0.193	-0.131 - 0.465	0.24
Arterial hypertension (No/Yes)	0.115	-0.092 - 0.387	0.22
Hyperlipidemia (No/Yes)	0.254	-0.215 - 0.638	0.31
Serum vitamin D	0.041	0.005 - 0.072	<0.05

**Table 3** Results of the multiple regression analysis considering age, gender, smoking habits, arterial hypertension, hyperlipidemia, and coronary artery calcium score as determinants of vitamin D level

Determinant	$\beta$	P-value	Multiple R <sup>2</sup>	F-test
Age (years)	0.07	0.42		
Gender (M/F)	0.09	0.13		
Smoking (No/Yes/Ex-smokers)	0.04	0.59		
Arterial hypertension (No/Yes)	-0.24	0.47		
Hyperlipidemia (No/Yes)	-0.18	<0.01	0.043	2.89
Coronary artery calcium score	-0.21	<0.001	0.231	1.54

## Discussion

Vitamin D deficiency poses a significant health challenge on a global scale, especially in developing nations.<sup>11</sup> In the current study, 52.1% of the total patients exhibited vitamin D deficiency (below 20 ng/ml). These figures align with the prevailing global prevalence.<sup>12</sup>

The assessment of vitamin D deficiency in various disorders, including CAD, has yielded conflicting results. Nevertheless, the presence of genetic polymorphisms in the vitamin D receptor suggests that differences related to ethnicity may influence the occurrence of vitamin D deficiency.<sup>13</sup> Our study revealed a discernible relationship between serum vitamin D levels and the severity of CAD. These findings diverge from those reported by *Ho et al.*<sup>14</sup> However, such discrepancies could be attributed to variations in geographical location, ethnicity, a lower CAD prevalence in their study, and the absence of female participants. Additionally, the relatively small sample size in our study may have contributed to these differences.

While previous global studies have consistently identified a connection between disorders related to coronary arteries and vitamin D deficiency, they have not directly delved into detailed angiographic data.<sup>15-21</sup> However, our current study reveals a noteworthy correlation between elevated CACS and the severity of vitamin D abnormalities (deficiency or insufficiency). Given the established role of CACS in predicting the plaque burden of coronary arteries, its significant association with low vitamin D levels in this study suggests a meaningful relationship between the severity of CAD and abnormal vitamin D levels.

In the evidence recently published by *Fang et al.*, genetically lowered 25(OH)D levels did not show an elevated risk of CAD.<sup>22</sup> This implies that the potential connection between vitamin D levels and CAD might be influenced by confounding factors or reverse causation. Vitamin D deficiency is linked to several conditions, including obesity, insulin resistance, diabetes, metabolic syndrome, hypertension, and hyperlipidemia.<sup>23-25</sup> Additionally, significant associations are observed between abnormalities associated with vitamin D and major

cardiovascular events, such as heart failure, myocardial infarction, sudden cardiac death, stroke, atrial fibrillation, and peripheral vascular disease.<sup>26,27</sup>

The precise understanding of the pathophysiology underlying connection between vitamin D deficiency and atherosclerosis is essential for grasping the underlying mechanisms and potential treatment implications. Numerous observational studies consistently reveal a reciprocal relationship between vitamin D deficiency and risk factors for atherosclerosis. Insufficient levels of vitamin D are linked to conditions like hypertension, dyslipidemia, insulin resistance, and obesity, all of which fuel the development and advancement of atherosclerosis.<sup>28</sup> Moreover, vitamin D deficiency correlates with indicators of hidden atherosclerosis, such as heightened carotid intima-media thickness and coronary artery calcification, underscoring its significant role in atherosclerosis progression. While the precise mechanisms linking vitamin D deficiency to atherosclerosis are still under investigation, one proposed mechanism revolves around the anti-inflammatory properties of vitamin D. It inhibits the production of pro-inflammatory cytokines and curtails the activation of immune cells involved in atherosclerotic processes. By modulating the inflammatory response, vitamin D may alleviate the inflammatory burden within arterial walls, consequently impeding atherosclerosis development and progression. Additionally, vitamin D might impact endothelial function, oxidative stress, and vascular smooth muscle cell proliferation—critical processes in atherosclerosis. By enhancing nitric oxide (NO) production, reducing oxidative stress, and dampening inflammation, vitamin D could ameliorate endothelial function. Furthermore, it may regulate the proliferation and migration of vascular smooth muscle cells, thereby contributing to plaque stability.<sup>29, 30</sup>

In conclusion, our findings, along with those of the majority of existing literature, suggest a correlation between vitamin D levels and the severity of CAD to some extent. Clearly, conducting additional well-designed studies with a larger cohort of Slovak patients is imperative to validate the role of vitamin D in both primary and secondary prevention of CAD.

## Limitations

While we established a significant association between vitamin D deficiency and CAD, certain limitations should be acknowledged. CAD is a chronic process typically initiated in childhood, yet we relied on a single laboratory report of vitamin D levels, which may not be indicative of past or lifelong vitamin D status. The fluctuation of vitamin D levels with seasons, though not assessed in our study, was mitigated by our sampling strategy, which covered all four seasons fairly evenly. Memory bias, inherent in relying on patient recollection regarding supplement avoidance during sampling, could not be entirely eliminated.

The experiment was conducted with a particular set of participants. Our study cohort consisted solely of patients residing in urban areas within the central region of Slovakia. The majority of these patients were undergoing treatment with statins. The study did not consider the dosage or duration of statin treatment, which is recognized to influence vitamin D levels.<sup>30</sup> Therefore, the findings cannot be generalized to other populations. Additionally, the study was observational and cross-sectional in nature, meaning it cannot establish causation but can only highlight statistical correlations. Finally, lack of a control group is one of the biggest limitations of the study which needs to be mentioned.

The known link between hypertension and vitamin D deficiency, coupled with the relationship between hypertension and CAD, introduces the possibility of confounding variables. Additionally, while CTA is a non-invasive imaging method with high sensitivity and specificity for identifying coronary plaques, the functional significance of these plaques remains elusive without diagnostic invasive selective coronary angiography. Nonetheless, ongoing advancements in CT devices with each new model are noteworthy. To clarify this hypothesis, further studies with larger samples, free from these limitations, are imperative.



## **Conclusions**

There was a correlation observed between vitamin D levels and both CACS and the severity of luminal stenosis caused by atherosclerotic plaques. Despite scientific societies' recommendations not endorsing vitamin D supplementation for preventing or slowing the progression of CAD, given the relatively high prevalence of cardiovascular disease in the study population, further research in this area is warranted. Such investigations could help evaluate the potential beneficial effects of such treatment in patients with these conditions.

Preprint

## References

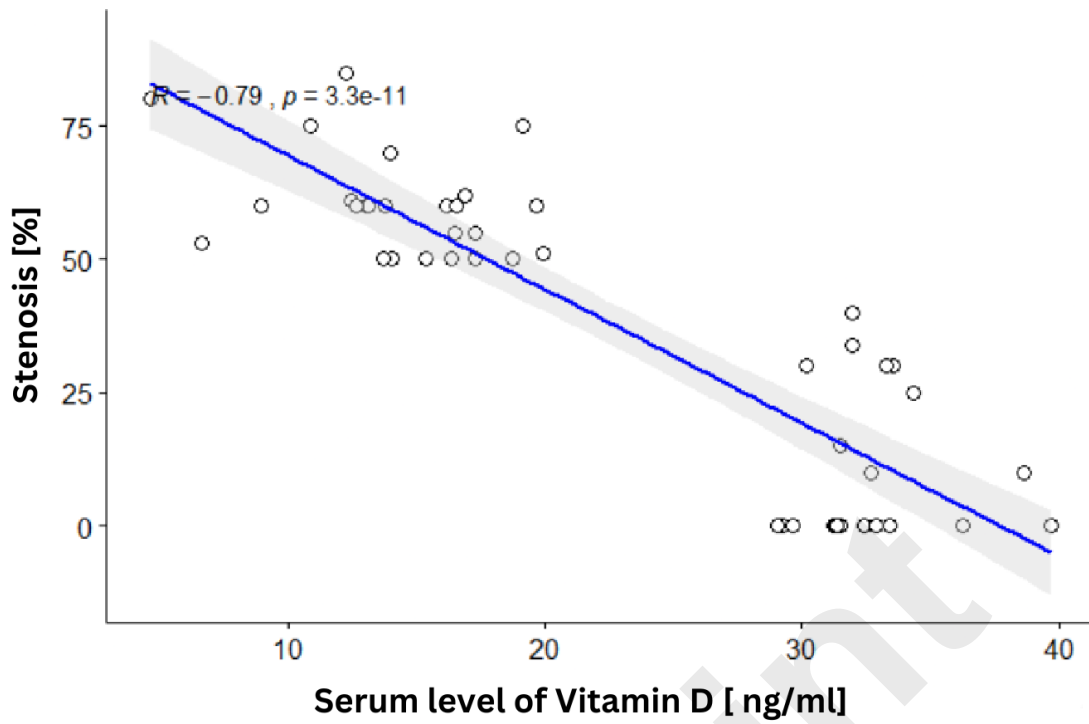
1. Saponaro F, Saba A, Zucchi R. An Update on Vitamin D Metabolism. *Int J Mol Sci.* 2020;21(18):6573.
2. Nakahara T, Dweck MR, Narula N, Pisapia D, Narula J, Strauss HW. Coronary Artery Calcification: From Mechanism to Molecular Imaging. *JACC Cardiovasc Imaging.* 2017;10(5):582-593.
3. Dabravolski SA, Markin AM, Andreeva ER, Eremin II, Orekhov AN, Melnichenko AA. Molecular Mechanisms Underlying Pathological and Therapeutic Roles of Pericytes in Atherosclerosis. *Int J Mol Sci.* 2022;23(19):11663.
4. Carbone F, Liberale L, Libby P, Montecucco F. Vitamin D in atherosclerosis and cardiovascular events. *Eur Heart J.* 2023;44(23):2078-2094.
5. Al-Oanzi ZH, Alenazy FO, Alhassan HH, et al. The Role of Vitamin D in Reducing the Risk of Metabolic Disturbances That Cause Cardiovascular Diseases. *J Cardiovasc Dev Dis.* 2023;10(5):209.
6. Verdoia M, Gioscia R, Nardin M, Rognoni A, De Luca G. Low levels of vitamin D and coronary artery disease: Is it time for therapy?. *Kardiol Pol.* 2022;80(4):409-416.
7. Manousaki D, Mokry LE, Ross S, Goltzman D, Richards JB. Mendelian Randomization Studies Do Not Support a Role for Vitamin D in Coronary Artery Disease. *Circ Cardiovasc Genet.* 2016;9(4):349-356.
8. Yan X, Wei Y, Wang D, et al. Four common vitamin D receptor polymorphisms and coronary artery disease susceptibility: A trial sequential analysis. *PLoS One.* 2022;17(10):e0275368.
9. Hanley DA, Cranney A, Jones G, Whiting SJ, Leslie WD; Guidelines Committee of the Scientific Advisory Council of Osteoporosis Canada. Vitamin D in adult health and disease: a review and guideline statement from Osteoporosis Canada (summary). *CMAJ.* 2010;182(12):1315-1319.
10. Hussain B, Mahmood A, Flynn MG, Alexander T. Coronary Artery Calcium Scoring in Asymptomatic Patients. *HCA Healthc J Med.* 2023;4(5):341-352.
11. Cui A, Zhang T, Xiao P, Fan Z, Wang H, Zhuang Y. Global and regional prevalence of vitamin D deficiency in population-based studies from 2000 to 2022: A pooled analysis of 7.9 million participants. *Front Nutr.* 2023;10:1070808.
12. Amrein K, Scherkl M, Hoffmann M, et al. Vitamin D deficiency 2.0: an update on the current status worldwide. *Eur J Clin Nutr.* 2020;74(11):1498-1513.
13. Fronczek M, Osadnik T, Banach M. Impact of vitamin D receptor polymorphisms in selected metabolic disorders. *Curr Opin Clin Nutr Metab Care.* 2023;26(4):316-322.

14. Ho JS, Cannaday JJ, Barlow CE, Reinhardt DB, Wade WA, Ellis JR. Low 25-OH vitamin D levels are not associated with coronary artery calcium or obstructive stenoses. *Coron Artery Dis.* 2015;26(6):521-525.
15. Pál É, Ungvári Z, Benyó Z, Várbíró S. Role of Vitamin D Deficiency in the Pathogenesis of Cardiovascular and Cerebrovascular Diseases. *Nutrients.* 2023;15(2):334.
16. Sabico S, Wani K, Grant WB, Al-Daghri NM. Improved HDL Cholesterol through Vitamin D Status Correction Substantially Lowers 10-Year Atherosclerotic Cardiovascular Disease Risk Score in Vitamin D-Deficient Arab Adults. *Nutrients.* 2023;15(3):551.
17. Dziedzic EA, Gąsior JS, Pawłowski M, Wodejko-Kucharska B, Saniewski T, Marcisz A, Dąbrowski MJ. Vitamin D level is associated with severity of coronary artery atherosclerosis and incidence of acute coronary syndromes in non-diabetic cardiac patients. *Arch Med Sci.* 2019;15(2):359-368.
18. Dziedzic EA, Przychodzeń S, Dąbrowski M. The effects of vitamin D on severity of coronary artery atherosclerosis and lipid profile of cardiac patients. *Arch Med Sci.* 2016;12(6):1199-1206.
19. Bakthavatchalam R, Bakthavatchalam S, Chandran I, et al. Association of Vitamin D with Risk Factors for Coronary Artery Disease. *Maedica (Bucur).* 2023;18(4):563-570.
20. Dziedzic EA, Gąsior JS, Saniewski T, Dąbrowski M. Vitamin D deficiency among Polish patients with angiographically confirmed coronary heart disease. *Pol Merkur Lekarski.* 2021;49(292):278-282.
21. El Mokadem M, Boshra H, Abd El Hady Y, Abd El Hameed AS. Relationship of serum vitamin D deficiency with coronary artery disease severity using multislice CT coronary angiography. *Clin Investig Arterioscler.* 2021;33(6):282-288.
22. Fang A, Zhao Y, Yang P, Zhang X, Giovannucci EL. Vitamin D and human health: evidence from Mendelian randomization studies. *Eur J Epidemiol.* Published online January 12, 2024.
23. Gariballa S, Al-Bluwi GSM, Yasin J. Frequency of Vitamin D Receptor Gene Polymorphisms in a Population with a very High Prevalence of Vitamin D Deficiency, Obesity, Diabetes and Hypertension. *Biomedicines.* 2023;11(4):1202.
24. Farahmand MA, Daneshzad E, Fung TT, et al. What is the impact of vitamin D supplementation on glycemic control in people with type-2 diabetes: a systematic review and meta-analysis of randomized controlled trails. *BMC Endocr Disord.* 2023;23(1):15.
25. Huang X, Yang Y, Jiang Y, Zhou Z, Zhang J. Association between vitamin D deficiency and lipid profiles in overweight and obese adults: a systematic review and meta-analysis. *BMC Public Health.* 2023;23(1):1653.

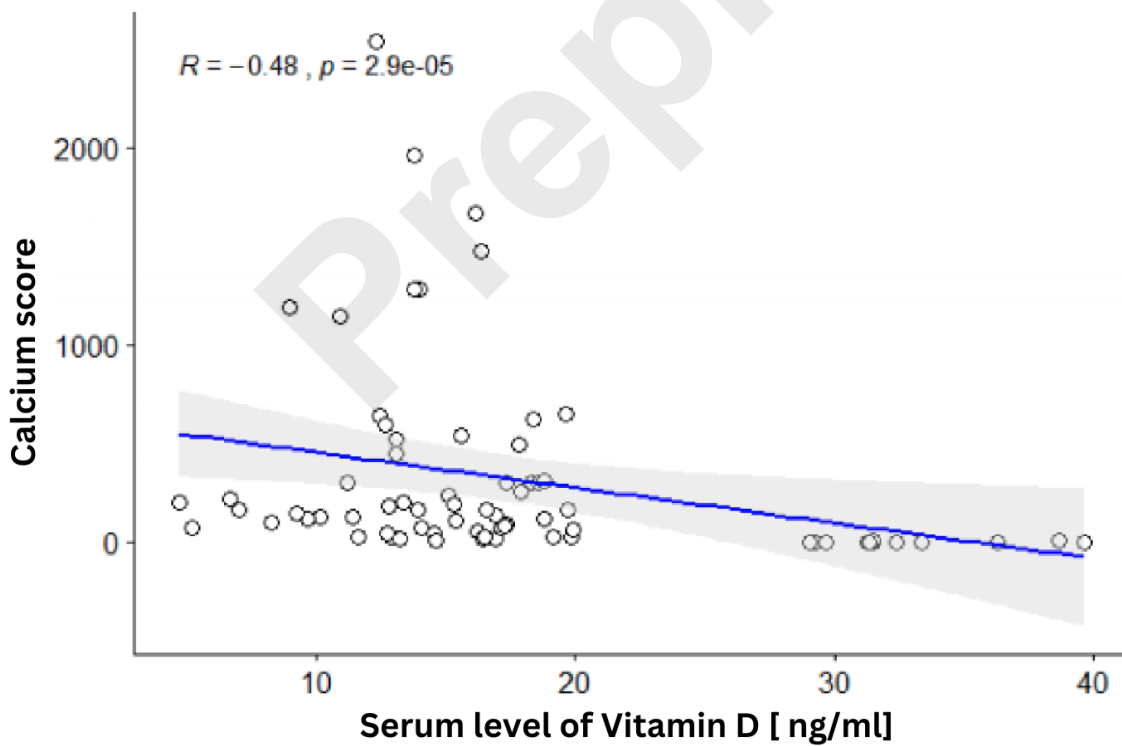
26. Zittermann A, Pilz S. Vitamin D and Cardiovascular Disease: An Update. *Anticancer Res.* 2019;39(9):4627-4635.
27. Knežević Praveček M, Vuković-Arar Ž, Miškić B, Hadžibegović I. Vitamin D Deficiency in Acute Coronary Syndrome - Clinically Relevant or Incidental Finding?. *Cent Eur J Public Health.* 2017;25(3):185-190.
28. Adeniyi OV, Masilela C, George JA. Prevalence of vitamin D deficiency and its association with cardiometabolic risk factors among healthcare workers in the Eastern Cape province, South Africa; cross-sectional study. *Sci Rep.* 2024;14(1):4756.
29. Roffe-Vazquez DN, Huerta-Delgado AS, Castillo EC, et al. Correlation of Vitamin D with Inflammatory Cytokines, Atherosclerotic Parameters, and Lifestyle Factors in the Setting of Heart Failure: A 12-Month Follow-Up Study. *Int J Mol Sci.* 2019;20(22):5811.
30. Dinca M, Serban MC, Sahebkar A, et al. Does vitamin D supplementation alter plasma adipokines concentrations? A systematic review and meta-analysis of randomized controlled trials. *Pharmacol Res.* 2016;107:360-371.

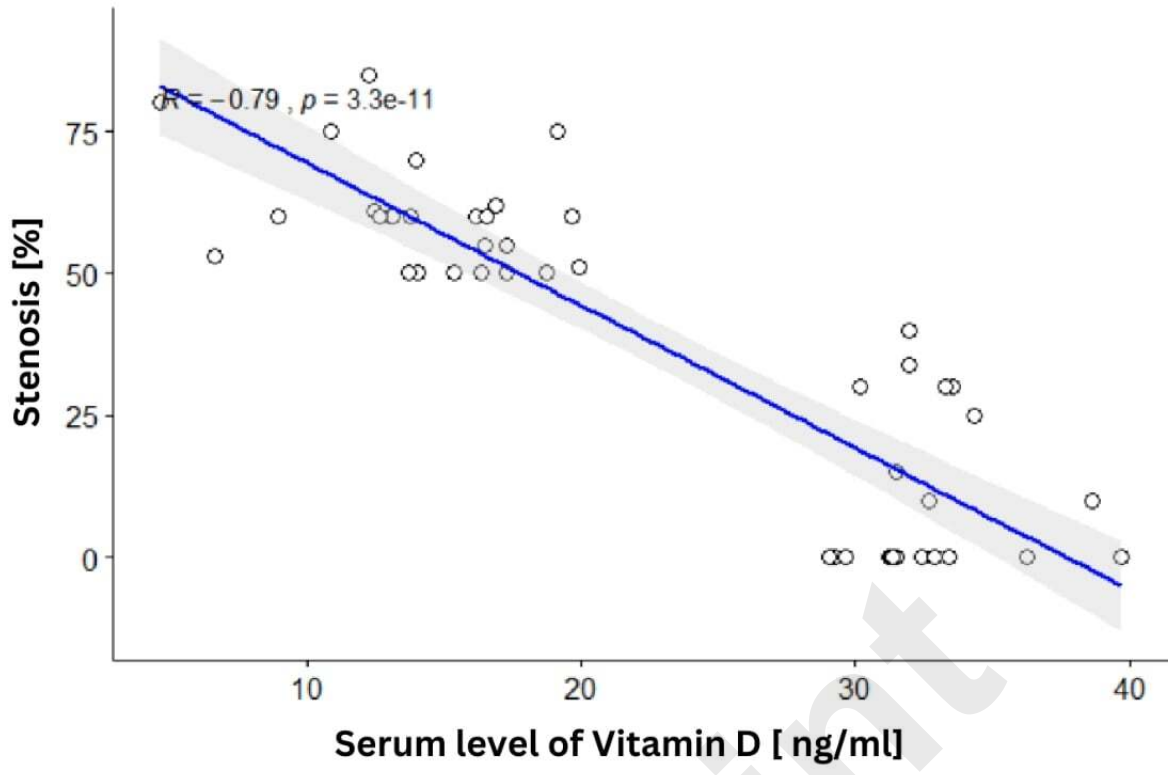
Preprint

**Figure 1** Spearman correlation between vitamin D levels in serum and the coronary artery stenosis



**Figure 2** Spearman correlation between vitamin D levels in serum and calcium score





Preprint