





Vitamin D and Its Metabolites Deficiency in Acute Coronary Syndrome Patients Undergoing Coronary Angiography: A Case–Control Study

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Background: Vitamin D deficiency is considered an emerging health problem that affects at least one billion patients worldwide. Calcitriol 1,25(OH)₂D₃ has several systemic effects, including anti-inflammatory, anti-thrombotic and anti-atherosclerotic impacts that explain its cardioprotective effects. The precise association between vitamin D and its metabolites and the value of supplements in acute coronary syndrome (ACS) is still controversial. This study aims to search the association between vitamin D₂, D₃, and metabolites and ACS in patients undergoing coronary angiography.

Materials and Methods: This was a case–control study on 73 consecutive adult patients with ACS undergoing coronary angiography compared to 50 controls without coronary artery disease and matched for age and sex from June 2019 till July 2019. Echocardiography and coronary angiography were done for all cases. Plasma vitamin D and its metabolites were measured at admission for all participants along with chemistry profiles.

Results: Vitamin D and its metabolites were statistically significantly lower in ACS patients than the controls. Multivariate regression analysis revealed that low levels of 25-hydroxyvitamin D (25(OH)D) and 1,25-dihydroxyvitamin D (1,25(OH)₂D) significantly predicted ACS occurrence; the other significant predictors were high systolic blood pressure (BP), high total cholesterol, and low high-density lipoprotein-cholesterol. Interestingly, vitamin D₂ and D₃ did not significantly predict ACS (p>0.05). We did not find a statistically significant association between the number of affected coronary vessels and vitamin D metabolites. Moreover, there was no statistically significant correlation between vitamin D and its metabolites and left ventricular ejection fraction measured by echocardiography.

Conclusion: There was a strong association between vitamin D and all its metabolites with ACS. Significantly, low 25(OH)D and 1,25(OH)₂D predicted ACS, but vitamin D₂ and D₃ did not. Large randomized controlled trials are needed to verify the beneficial values of vitamin D supplementation in ACS patients.

Keywords: vitamin D₃, vitamin D₂, 25-hydroxyvitamin D, 1,25-dihydroxyvitamin D₃, acute coronary syndrome, coronary artery disease

Introduction

Coronary artery disease (CAD) is considered the leading cause of morbidity and mortality worldwide.¹ New emerging risk factors have been proposed to be significant predictors of atherosclerosis and subsequent complications. Among them, vitamin D deficiency may be involved in the pathogenesis of acute coronary syndrome (ACS).²

The prevalence of vitamin D deficiency has increased among all populations and ethnicities in the last few years.³ It was identified as a pandemic affecting approximately one-third of the Americans,⁴ 60% of the Canadians,⁵ and 40% of the Europeans.⁶ Additionally, in Saudi Arabia, low 25-hydroxy vitamin D (25(OH)D) was found in 36.8% to 62.65% of females and 17.7% to 40.6% of males.^{7,8}

In addition to the effects of vitamin D on calcium homeostasis,⁹ it has beneficial anti-inflammatory and anti-atherosclerotic effects.¹⁰ Moreover, hypovitaminosis D has been associated with atherosclerotic cardiovascular disease (CVD),² higher post-ACS complications,¹¹ and metabolic diseases, eg, metabolic syndrome and insulin resistance.¹²

While the exact mechanism of increased cardiovascular risk in patients with vitamin D deficiency has not yet been established, multiple hypotheses have been postulated. Vitamin D receptors (VDRs) have been distributed widely throughout the cardiovascular system.¹³ Vitamin D reduces cardiac ischemia-reperfusion injury and reactive oxygen species¹⁴ and shows favourable effects on inflammation and thrombosis.¹⁵ Additionally, low 25(OH)D levels promote atherosclerosis, chronic inflammation, endothelial dysfunction and arterial calcification.¹⁶ Moreover, vitamin D insufficiency may activate the renin-angiotensin system¹⁷ and increase insulin resistance, endothelial dysfunction, inflammation, platelet function, and blood pressure (BP).¹⁸ Furthermore, an experimental trial revealed that vitamin D supplementation suppressed vascular inflammation by inhibiting the Nuclear Factor- κ B (NF- κ B) pathways and decreasing the process of atherosclerosis and hence subsequent CAD.¹⁹ Additionally, a study demonstrated an inverse association between matrix metalloproteinase 9 and vitamin D in 139 acute myocardial infarction (AMI) patients.²⁰

The two forms of vitamin D are vitamin D₂ (D₂), which is present in plant foods, and vitamin D₃ (D₃), which is formed in the skin upon exposure to sunlight and from animal food.²¹ D₃ is three times more effective than D₂ in humans.²² Both D₂ and D₃ undergo hepatic hydroxylation at position 25 to generate 25(OH)D, ie, calcidiol.²³ Further, one α -hydroxylation is taken place to produce 1,25-dihydroxyvitamin D (1,25 (OH)₂D), ie, calcitriol, which is the active form of vitamin D, but it circulates in lower concentrations when compared to 25 (OH)D.²⁴

Although several studies were done on the 25(OH)D, there is no single study on the association of different

forms of vitamin D and their metabolites and the acute coronary syndrome on our population. Therefore, it would help to study the potential association between vitamin D forms with their metabolites and ACS in our population.

Methodology

Study Population

The present study is a case–control study that included 73 consecutive adult patients with ACS admitted to the coronary care unit at the Madinah Cardiac Centre, a tertiary care hospital, Al-Madinah Al-Munawara, that is considered one of the referral centres for the western region of Saudi Arabia, during the period from June 2019 till July 2019. The controls were 50 age- and gender-matched subjects free from previous CVD recruited from the attendants or relatives of noncardiac patients. We included adult patients (age >18 years old) with ACS. We excluded patients with chronic coronary syndrome, pregnancy, malignancy, thyroid disease, advanced chronic kidney disease (stage 4 or 5), and liver disease.

Collection of Data

Clinical Data

We interviewed the patients and obtained a full history and complete examination. Specifically, the history included personal demographic data and the traditional CVD risk factors. Furthermore, a complete cardiac examination, electrocardiogram (ECG), echocardiography, and diagnostic coronary angiography were done on all patients.

Criteria of Diagnosis of Risk Factors

Hypertension was considered if the patient was previously diagnosed or on active anti-hypertensive treatment.²⁵ The patient was considered diabetic if glycated haemoglobin (HbA1c) is 6.5% or more, previously diagnosed, or on antidiabetic treatment.²⁶ Smoking was considered if patients smoked at least 100 cigarettes or currently smoking. While Ex-smokers were patients who smoked 100 cigarettes or more, but they are not currently smoking, and they have not smoked for two years. Ex-smokers and never-smokers were considered non-smokers.²⁷ Serum cholesterol and its lipoproteins are directly linked to the development of CVD. General population studies suggest that the optimal levels of total cholesterol and LDL-C is 150 mg/dL (3.8 mmol/L), and 100 mg/dL (2.6 mmol/L), respectively.²⁸ In cases with a very high risk of CVD, ie, secondary prevention, the target is a reduction of LDL-C

$\geq 50\%$ from baseline and an LDL-C goal of < 1.4 mmol/L (< 55 mg/dL).²⁹

HDL-C is associated with the risk of CVD³⁰ even though no specific goals have been set for HDL-C in clinical trials.²⁹

Diagnosis of ACS

The acute myocardial infarction requires at least two of the following: symptoms suggestive of ischemia, diagnostic ECG, and elevated cardiac biomarkers.³¹ ECG suggestive of ACS includes presumably new pathologic Q waves or a new left bundle branch block. Moreover, ST-elevation is ≥ 0.1 mV in 2 or more contiguous limb leads and/or ≥ 0.2 mV ST elevation in chest leads.³² Unstable angina was diagnosed in patients presenting with symptoms suggestive of ischemia without cardiac biomarker elevation, and ECG might indicate ischemia.³³

Laboratory Data

Blood samples were collected from the study subjects under aseptic measures by venipuncture within 24 h of admission. The samples were immediately centrifuged on-site at $2000 \times g$ for 10 minutes at 4°C and then stored at -70°C until the analysis. Serum samples were thawed at room temperature just before analysis. Serum samples were analyzed at the College of pharmacy-Taibah University laboratories. Serum levels of vitamin D, including D_2 , D_3 , $25(\text{OH})\text{D}$, and $1,25(\text{OH})_2\text{D}$, were measured by the ultra-performance liquid chromatography (UPLC) system with a tandem mass detector system,³⁴ which was made up of Waters Quattro Premier XE tandem mass spectrometer (Waters Corporation, USA).

Regarding the diagnosis and categorization of serum levels of $25(\text{OH})\text{D}$ deficiency that was based on the United States Endocrine Society guideline, $25(\text{OH})\text{D}$ deficiency was serum level ≤ 20 ng/mL, and $25(\text{OH})\text{D}$ insufficiency with serum level between 20 and 29.99 ng/mL, while the normal values were ≥ 30 ng/mL.³⁵

Ethical Considerations

This work was done and accepted by the research ethics committee of the faculty of Medicine, Taibah University, Al-Madinah Al-Munawara, Saudi Arabia. All patients who were enrolled in the study agreed on written informed consent. Patients were told that their details would be confidential. The current study was conducted in accordance with the World Medical Association Declaration of Helsinki.

Statistical Analysis

SPSS 21.0 software was used for data analysis (SPSS Inc. Chicago, USA). All quantitative data were presented by the mean and standard deviation (SD), and they were compared by Student's *t*-test when data were normally distributed, while the Mann–Whitney test has been used when data were not normally distributed. Additionally, all qualitative data were expressed as frequency and per cent, and they were compared by Chi-square test. Pearson's correlation was used to determine the correlation between two quantitative variables with normally distributed data. Meanwhile, Spearman correlation was used to test the correlation between two quantitative variables having non-normally distributed data. Moreover, multiple logistic regression models (the enter mode) have been used to give an adjusted odds ratio and 95% confidence interval of the effect of different risk factors for ACS. A statistical significance was considered when the *p*-value was below 0.05.

Results

The current study included 73 cases with ACS compared to 50 volunteers without a history of ACS as a control group. Overall, the participants' age ranged from 30 to 83 years old, with an overall mean \pm SD of 56.7 ± 11.5 . The baseline characteristics of our study population were reported in Table 1. In this study, statistically significant differences have been found between cases and controls regarding the medical history of diabetes and hypertension ($p < 0.001$) (Table 1).

In Table 2, there were statistically significant differences between the ACS cases and the controls regarding the systolic blood pressure ($p = 0.002$). In addition, blood glucose was significantly higher in the ACS than in the controls. Notably, lipid profile results were also significantly different between the two groups ($p < 0.001$). Haematological parameters such as haemoglobin, white blood cell count, and platelet count showed a significant difference between the ACS and the control groups ($p < 0.001$).

To assess the possible association between vitamin D and the development of ACS, we measured vitamin D_2 , D_3 and the metabolites and found that the mean \pm SD of all measured vitamin D forms and the metabolites were significantly lower in the ACS group than in the control group with a *p*-value < 0.001 (Table 3). In addition, the

Table 1 General Characteristics of Study Participants

Variables	Participants (n=123)				p-value ^a
	ACS Cases* (n=73)		Control (n=50)		
	No	%	No	%	
Age:					
<45	6	8.2	9	18.0	0.103
≥45	67	91.8	41	82.0	
Gender:					
Male	57	78.1	38	76.0	0.78
Female	16	21.9	12	24.0	
History of Diabetes:					
No	27	37.0	42	84.0	<0.001*
Yes	46	63.0	8	16.0	
History of hypertension:					
No	31	42.5	41	82.0	<0.001*
Yes	42	57.5	9	18.0	
Smoking:					
No	47	64.4	37	74.0	0.26
Yes	26	35.6	13	26.0	
BMI:					
Underweight or normal	26	35.6	16	32.0	0.67
Overweight or obese	47	64.4	34	68.0	

Notes: ^aChi square test. *P-value < 0.05 is considered statistically significant.

Abbreviations: ACS, acute coronary syndrome; BMI, body mass index; no, number.

study showed that 25(OH)D was either deficient or insufficient in all ACS cases (Figure 1).

In the current study, all variables significantly ($p < 0.05$) associated with ACS in univariate analysis were included

in multivariate regression analysis models in addition to age and gender. High systolic blood pressure and high total cholesterol level were independently associated with ACS among participants (OR=1.128, 95% CI=1.007–1.264) and (OR=8.286, 95% CI=1.073–63.972), respectively. While the lower levels of high-density lipoprotein cholesterol (HDL-C), 25(OH)D, and 1,25(OH)₂D were independently associated with ACS among participants (OR=0.001, 95% CI=<0.001–0.040; OR=0.218, 95% CI= 0.065–0.732; OR=0.545, 95% CI=0.311–0.956), respectively (Table 4).

As shown in Figure 2, more than half of the ACS cases (65.8%) presented with single-vessel CAD. While the frequencies of different affected coronary arteries; left main (LM), left anterior descending (LAD), left circumflex (LCX), and right coronary artery (RCA), were 11.0%, 54.8%, 34.2%, and 47.9%, respectively among the studied ACS cases. Moreover, the mean±SD of the left ventricular ejection fraction was 36.59±12.46 in the ACS group. Interestingly, no significant correlations ($p > 0.05$) have been found between all the vitamin forms and metabolites and the number of affected vessels (it was ranging from 1 to 3 vessels) and the left ventricular ejection fraction among the studied group of ACS cases (Table 5). Meanwhile, a statistically significant correlation has been found between 25-hydroxyvitamin D and 1,25-dihydroxyvitamin D ($p < 0.001$, $r = 0.88$) among the same group of patients (Figure 3).

Discussion

Deficiency of vitamin D is a worldwide health problem with a high prevalence among the general population.³⁶ Most

Table 2 Clinical Parameters, Blood Glucose, Lipid Profile, and Hematological Tests Among Participants

Parameters	Participants (n=123)		P-value
	ACS Cases (n=73) Mean±SD	Controls (n=50) Mean±SD	
Systolic BP (mmHg)	129.60±21.72	118.48±13.29	0.002**
Diastolic BP (mmHg)	75.42±10.64	77.54±10.29	0.275**
Glucose Level (mg/dL)	168.86±63.39	125.75±25.58	<0.001**
TC (mmol/L)	4.48±1.41	3.05±0.44	<0.001**
LDL-C (mmol/L)	2.90±1.24	1.92±0.69	<0.001*
HDL-C (mmol/L)	1.00±0.25	1.18±0.12	<0.001**
Triglycerides (mmol/L)	1.24±0.77	1.21±0.81	0.361*
Hb (g/dL)	13.21±1.99	11.25±0.89	<0.001**
RBCs (10⁶/uL)	4.63±0.63	4.69±0.64	0.653**
Total WBCs (10³/uL)	10.55±3.29	7.18±1.29	<0.001**
Platelets (10³/uL)	267.98±75.36	215.10±25.37	<0.001**

Notes: *P-value of Mann Whitney test. **P-value of Student's t-test. P-value < 0.05 is considered statistically significant, and is bolded.

Abbreviations: ACS, acute coronary syndrome; BP, blood pressure; TC, total cholesterol; LDL-C, low-density lipoproteins cholesterol; HDL-C, high-density lipoproteins cholesterol; Hb, haemoglobin; RBC, red blood cells; WBC, white blood cells.

Table 3 Serum Levels of Vitamin D₂, D₃, and the Metabolites Among Participants

Parameters	Participants (n=123)		P-value
	ACS Cases* (n=73) Mean±SD	Controls (n=50) Mean±SD	
Vitamin D ₂ (ng/mL)	23.48±5.12	31.64±6.26	<0.001*
Vitamin D ₃ (ng/mL)	23.13±5.64	32.35±6.31	<0.001*
25(OH)D (ng/mL)	14.65±3.46	24.42±4.25	<0.001*
1,25(OH) ₂ D (pg/mL)	45.56±8.85	65.15±7.15	<0.001*

Notes: Student's t-test was used. *P-value < 0.05 is considered statistically significant, and is bolded.

Abbreviations: ACS, acute coronary syndrome; 25(OH) D, 25-hydroxyvitamin D; 1,25(OH)₂D, 1,25-dihydroxyvitamin D.

studies were conducted on the association between 25(OH) D and myocardial infarction. The current study was a single-centre study conducted in Saudi Arabia, compared serum vitamin D₂, D₃, their metabolites in ACS patients, and age and gender-matched controls without CVD.

Our main finding was a highly significant difference between the ACS cases and the controls regarding all forms of vitamin D and the two metabolites. The mean vitamin D₂, D₃, 25(OH)D, and 1,25(OH)₂D were significantly lower in the ACS cases compared to the controls (Table 3). Moreover, the 25(OH)D deficiency was significantly prevalent in our cohort, and there was a significantly higher prevalence in ACS patients (91.4%) compared to the control group (14%).

The regression analysis demonstrated the strongly associated predictors with ACS. Low serum levels of 25(OH)D and 1,25(OH)₂D significantly predicted ACS occurrence. Moreover, high total cholesterol (TC) and HDL-C significantly predicted ACS, as did high systolic BP. Conversely, the D₂ and D₃ did not significantly predict the occurrence of ACS.

In line with our findings, several studies have shown the association between deficiency of 25-hydroxyvitamin D and the risk of CAD or AMI in multiple ethnicities.^{37–40} Moreover, 25(OH)D deficiency has been associated with more severe CAD on coronary angiography.⁴¹ While subjects with normal serum levels of 25(OH)D were at a lower risk for AMI compared to 25(OH)D deficient ones after adjustment for potential confounders.⁴²

Additionally, several studies showed that low 25(OH)D prevalence was as high as 68–96% in ACS patients.^{43–46} Of note, the median serum 25(OH)D was as low as 18 ng/mL in 206 AMI patients.⁴⁷

On the other side, the abnormally high 25(OH)D can be associated with high major adverse cardiac events and total mortality, that is, a U-shape curve.⁴⁸ Rajasree et al enrolled 143 men with CAD and 70 male controls. High levels of 25(OH)D (>89ng/mL) were found in 59.4% of the CAD cases vs 40.6% in the controls. CAD patients with high serum 25(OH)D levels had more than a three-fold increase likelihood of having CAD, even after adjusting with multivariate logistic regression.⁴⁹

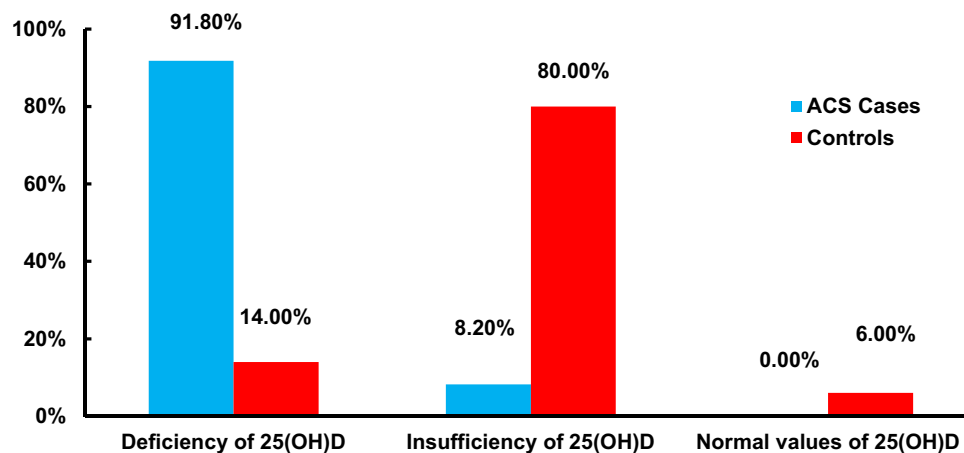


Figure 1 Distribution of serum 25(OH)D among the study participants. 25(OH)D deficiency: serum level ≤20ng/mL, insufficiency: 20.01–29.99ng/mL, and normal values: >30ng/mL.

Abbreviations: ACS, acute coronary syndrome; 25(OH)D, 25-hydroxyvitamin D.

Table 4 Factors Associated with ACS Among the Study Participants

Variables	P-value	Adjusted (OR)**	95% CI
25(OH)D (ng/mL)	0.014*	0.218	0.065–0.732
1,25(OH)₂D (pg/mL)	0.034*	0.545	0.311–0.956
Systolic BP	0.038*	1.128	1.007–1.264
TC	0.043*	8.286	1.073–63.972
HDL-C	0.011*	0.001	<0.001–0.040

Notes: *P-value < 0.05 is considered statistically significant, and is bolded. **Odds ratio has been adjusted by regression analysis for all possible confounders.

Abbreviations: ACS, acute coronary syndrome; 25(OH) D, 25-hydroxyvitamin D; 1,25(OH)₂D, 1,25-dihydroxyvitamin D; BP, blood pressure; TC, total cholesterol; HDL-C, high-density lipoproteins cholesterol.

Regarding 1,25(OH)₂D or calcitriol status, we found that the active vitamin D metabolite calcitriol is directly correlated to calcidiol. This finding agreed with the work of Dobing et al,⁵⁰ as they showed a significant association between low calcidiol and calcitriol and cardiovascular and total mortality. Renal function is the main factor affecting the serum level of calcitriol.⁵¹ Renal impairment causes a defective renal hydroxylation of calcidiol; therefore, it was recommended to supplement calcitriol or one of its analogues to those patients to raise their blood level, resulting in increased survival.⁵² Notably, the active form of vitamin D has both cardioprotective and pleiotropic effects and increases thrombomodulin and decreases tissue factor.⁵³

In line with our results on the systolic BP as a predictor of ACS, “The Third National Health and Nutrition Examination Survey” has reported a significant increase in the prevalence of hypertension [OR = 1.30, p < 0.001] in patients with low 25(OH)D (<21 ng/mL) compared to those patients with level more than 37 ng/mL.⁵⁴ In addition, a study confirmed the relationship between the TC, low-density lipoprotein cholesterol (LDL-C), HDL-C, and non-HDL-C and ACS vs control group.⁵⁵

Concerning LVEF by echocardiography, we did not find a statistical correlation between the LVEF and vitamin D₂,

D₃, and metabolites [calcidiol and calcitriol]. In agreement with our finding, Pandit et al⁵⁶ found no significant association between LVEF and 25(OH)D despite adjustment by age and hypertension treatment. Conversely, Pilz et al⁵⁷ have shown that vitamin D decreased with impaired LV function. Moreover, vitamin D deficiency was significantly associated with sudden cardiac death and death due to heart failure. This study was done on 3,299 patients who were referred for coronary angiography. They did not recruit ACS patients. Our study was on ACS patients and recruited a relatively small number of cases. Interestingly, the early LVEF few days after AMI did not predict the subsequent value. As demonstrated with the study of Chew et al in 2018, they studied 3 cohorts with AMI and found that LVEF showed a 25% to 37.5% increase from the baseline EF during the presentation of AMI to 2–8 weeks post-AMI.⁵⁸

We did not find a statistically significant difference in the vitamin D level between one-vessel and three-vessel disease regarding the coronary angiography results. On the contrary, Syal et al found a significantly higher incidence of double- and triple-vessel CAD in patients with low 25(OH)D compared to those with a normal 25(OH)D level (53% and 38%, respectively).⁴¹ Of note, we do not have any ACS patients with an optimal range of 25(OH)D. Thus, all our patients were in the deficient and insufficient categories of 25(OH)D.

Though the deficiency of vitamin D was significantly associated with CAD and ACS in many studies, the role of vitamin D supplementation was investigated recently. In the large randomized trial for primary prevention, the Vitamin D and Omega-3 Trial (VITAL), a relatively large supplemental vitamin D (2000 IU) dose failed to reduce the composite endpoint of major cardiovascular events.⁵⁹

Conclusions

This study showed that vitamin D and all its metabolites were statistically significantly deficient in all patients with

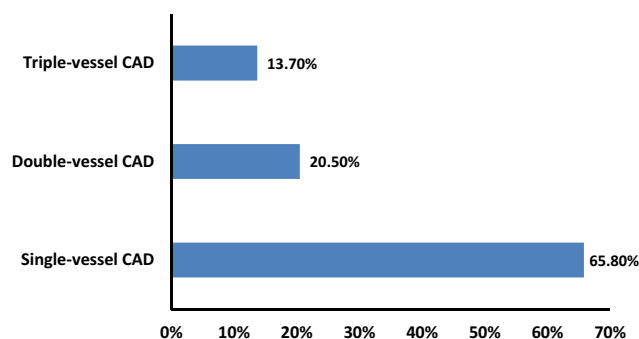


Figure 2 Percentage of single, double and triple vessel CAD among the studied cases of ACS (n=73).

Abbreviations: ACS, acute coronary syndrome; CAD, coronary artery disease.

Table 5 Correlations of Vitamin D₂, D₃, and the Metabolites with the Number of Affected Coronary Arteries and LV Ejection Fraction Among ACS Cases (n=73)

Parameters	Number of Affected Coronary Arteries (1–3)		LVEF (%)	
	r	p-value*	r	p-value*
Vitamin D ₂ (ng/mL)	0.075	0.529	0.087	0.467
Vitamin D ₃ (ng/mL)	0.035	0.771	0.161	0.327
25(OH)D (ng/mL)	0.044	0.710	0.155	0.190
1,25(OH) ₂ D (pg/mL)	0.023	0.846	0.113	0.339

Notes: Spearman correlation was used. *P-value < 0.05 is considered statistically significant.

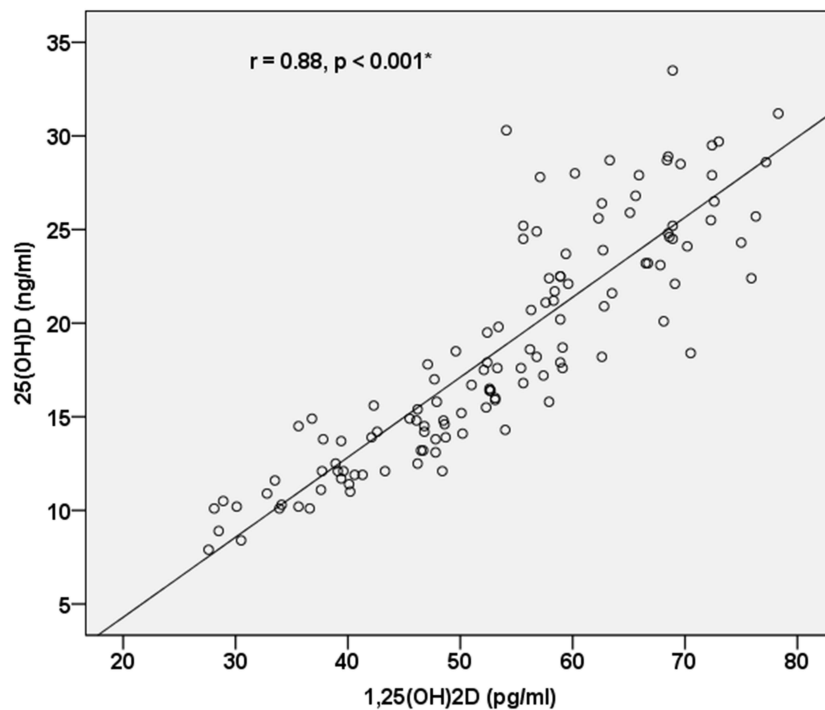
Abbreviations: ACS, acute coronary syndrome; r, Spearman correlation coefficient; LVEF, left ventricular ejection fraction; 25(OH) D, 25-hydroxyvitamin D; 1,25(OH)₂D, 1,25-dihydroxyvitamin D.

ACS compared with the controls. In addition, we demonstrated that 25(OH)D, 1,25(OH)₂D, high systolic BP, total cholesterol, and low HDL-C were statistically significant ACS predictors. Conversely, there was no association between the number of affected coronary vessels and vitamin D and its metabolites.

Future Research Perspectives

The clinical benefits of screening and treating vitamin D deficiency might reduce CVD morbidity and mortality. Thus, vitamin D supplementation is recognized as a potential therapeutic agent in patients with vitamin

D insufficiency/deficiency and cardiovascular diseases. However, vitamin D administration has many challenges, including possible high toxicity levels and low bioavailability. Thus, a potential new method such as using nanotechnologies to deliver the optimal amount of vitamin D either by incorporating into foods or therapies to enhance vitamin D bioavailability and effectiveness and overcome some of the drawbacks of the traditional way of administration.⁶⁰ We suggest that acute administration of 25(OH)D, or 1,25(OH)₂D for admitted ACS patients in large interventional randomized clinically controlled trials might improve patients' outcomes. However, further

**Figure 3** Correlation between 25(OH) D and 1,25(OH)₂D levels among the studied acute coronary syndrome cases (n=73).

Notes: Pearson's correlation was used. *P-value < 0.05 is considered statistically significant, and bolded.

Abbreviations: ACS, acute coronary syndrome; r, Pearson's correlation coefficient; 25(OH)D, 5-hydroxyvitamin D; 1,25(OH)₂D, 1,25-dihydroxyvitamin D.

factors should be considered with supplementation of vitamin D. Those include but are not limited to parathormone, phosphate, renin, and fibroblast growth factor.

Dual actions of VDR in transcription and non-genomic activation of RhoA–Rho-associated coiled kinase (ROCK), p38 mitogen-activated protein kinase (p38MAPK), and mitogen- and stress-activated kinase 1 (MSK1) may be a useful molecular therapeutic target to develop novel cardioprotective drugs.⁶¹ It was reported that vitamin D mitigated cardiomyocyte apoptosis, mitochondrial fission, mitophagy, and their ultrastructural abnormalities. Targeting signalling networks of inflammation and oxidative stress may be helpful. However, multitargeting cardioprotective therapeutic strategies should be adopted to tackle both inflammatory and non-inflammatory cardiac injury pathways consequent to ACS. Bio-intervention in complex structure one is suggested as a new therapeutic target. Mitophagy has dual beneficial and harmful effects for mitochondrial control, quality and cell survival. There are several creative multicenter approaches to hasten translational research for developing novel molecular therapy for cardioprotection and prevention of ACS. These include targeting mitochondrial damage, oxidative stress, and calcium overload, obstructed coronary microcirculation, activation of platelets and other critical pathways. Another innovative targeting includes engineered exosomal peptides that enhanced cardiomyocyte uptake, reduced apoptosis and high cardiac protection. Experimental intramyocardial microinjection opens a new arena for future gene therapy and improves the functional capacity of the heart. So, advances in nano-cardio medicine have huge potential future promising novel therapeutic agents for diagnosing, treating, and preventing acute coronary syndromes.

Strength and Limitations

Several studies were carried out on the association between the 25(OH)D levels and the ACS,^{37,39} yet, there is no single study, up to the best of our knowledge, on the association of different forms and metabolites of vitamin D and the occurrence of ACS on our population. Therefore, the strength of the current research is the study of the potential association between different forms and metabolites of vitamin D and ACS in our population. Our study has some limitations that need to be acknowledged. The sample of patients was only conducted from a single centre at a single time point. A small study sample reduces the power of event-free survival analysis, and the results obtained thus cannot be generalized globally to all ACS patients. These data can only be

used to generate hypotheses that can be used in the future for a confirmatory study, ie, a mega randomized clinical trial. Since this, an observational study, many different techniques that can be applied to prevent or control for confounding could not be used. The most commonly confounding factors that are hard to measure and adjust are limited physical activity, sun exposure, and diet. Moreover, the study did not consider the socioeconomic variables between the two groups that significantly influence vitamin D deficiency risk.

Abbreviations

ACS, acute coronary syndrome; AMI, acute myocardial infarction; BP, blood pressure; BMI, body mass index; CAD, coronary artery disease; CVD, cardiovascular disease; DM, diabetes mellitus; D2, vitamin D2; D3, vitamin D3; ECG, electrocardiogram; HbA1c, glycated haemoglobin; HDL-C, high-density lipoprotein cholesterol; LDL-C, low-density lipoprotein cholesterol; LVEF, left ventricular ejection fraction; NF- κ B, nuclear factor- κ B; TC, total cholesterol; VDRs, vitamin D receptors; VITAL, vitamin D and Omega-3 trial; 25(OH)D, 25-hydroxyvitamin D; 1,25(OH)₂D, 1,25-dihydroxyvitamin D; ROCK, RhoA–Rho-associated coiled kinase; p38MAPK, p38 mitogen-activated protein kinase; MSK1, mitogen- and stress-activated kinase 1.

Disclosure

The authors report no conflicts of interest related to this work.

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