



Examine the Vitamin-D Status of HCV-Infected Patients and Control in Lahore, Pakistan

**Muhammad Abbas ^{a*}, Hafiz Usman Ghani ^b, Huda Rauf ^c, Saba Zulfiqar ^d,
Muhammad Usman ^e and Abdul Rehman ^e**

^a Hebei North University, China.

^b Hebei Medical University, China.

^c DHQ Teaching Hospital, Sargodha, Pakistan.

^d Department of Biochemistry, University Medical and Dental College, Faisalabad, Pakistan.

^e Pakistan Institute of Medical Sciences, Islamabad, Pakistan.

Authors' contributions

This work was carried out in collaboration among all authors. All authors read and approved the final manuscript.

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ABSTRACT

Introduction: Hepatitis C virus is a serious health issue that can remain fatal owing to their dire repercussions. HCV causes approximately 710,500 deaths each year, has over 190 million individuals affected

Objective: Serum Vitamin-D has an important part in inflammatory and infections illnesses, particularly liver diseases. The purpose of this experiment was to examine the Vitamin-D status of HCV-diseased cases and control in Lahore, Pakistan.

Methods: From the 25th of August, 2019 to the 25th of February, 2020, we conducted randomized cross-sectional research on 78 people at Services Hospital in Lahore. Fifty-one individuals tested positive for hepatitis C RNA-PCR (25 salaried cirrhotic and 25 decompensated cirrhotic cases). In addition, 24 healthy controls were selected who did not have liver problems. ARTUS ® HCV QS-RGQ V1 was used to perform HCV RNA-PCR. Chemiluminescence was used to determine vitamin D levels. For descriptive statistics, SPSS version 20 was employed.

Results: When compared to untreated (32.43 ng/mL), average level of Vitamin-D in Treated patients remained considerably inferior in supplied and fluid resuscitation hepatocellular carcinoma

(28.88 ng/mL & 21.67 ng/mL, respectively). In this research, 78.6 percent of HCV patients had suboptimal Vitamin-D levels. Vitamin-D deficiency (22-28 ng/mL) was seen in both healthy people (46.9 percent) and HCV sufferers (38.3 percent) (P 0.002). Furthermore, Vitamin-D levels were shown to have an inverse association with more multi - organ failure, having 56.3 percent of respiratory failure cirrhosis people undergoing from Vitamin-D insufficiency, relative to 14.7 percent in the recovered cirrhotic groups (P 0.0002).

Conclusion: Suboptimal levels of Vitamin-D (inadequacy before lack) remain more common in hepatitis C comparison to well controls. Vitamin-D insufficiency were found to be directly related to somatic symptoms.

Keywords: Serum Vitamin-D; inflammatory and infections; illnesses; HCV-infected.

1. INTRODUCTION

Hepatitis C virus is a serious health issue that can remain fatal owing to their dire repercussions. HCV causes approximately 710,500 deaths each year, has over 190 million individuals affected [1]. HCV frequency is substantial in Pakistan, and it contributes significantly to the burden of liver disease. In reality, it is a Endemic of epic proportions, with the one in every 25 Pakistanis afflicted with HCV. As a result, as a national priority, efforts for management and cure of such a Endemic are required [2]. It seems to be well documented that vitamin D inadequacy is linked to inflammatory and pathogenic disorders, especially liver illnesses standing out. Vitamin-D blood levels have an opposite relation with the degree of liver problems and the progression of liver fibrosis [3]. Several studies have found a link between 25-hydroxy Vitamin D levels in blood also hepatitis C. Ultraviolet B (UVB) irradiation induces the production of up to 92 percent of Vitamin-D (in the form of cholecalciferol - D3) in the skin. The remaining is obtained orally through foods and supplements. Fish, eggs, and milk are deemed sources of D3 in the diet. Vitamin-D2 (ergocalciferol) is sourced from plants and is not generated in humans [4]. Vitamins D3 and D2 are both usually harmless. Vitamin-D activation happens in two phases hydroxylation reaction; the first hydroxylation at the 26th carbon atom occurs in the liver, which is one of causes wherefore Vitamin-D insufficiency is implicated in the pathogenesis liver illness. In the convoluted tubule of the kidneys, the first carbon of calcitriol is hydroxylated. This results in the production of completely activated Vitamin-D as 1, 26(OH) 2 D3, popularly recognized as calcitriol [5].

Vitamin-D levels are estimated by testing blood concentrations from both forms of 25-(OH) D3 and D2, although assessing serum calcitriol (activation form) is difficult given the short life span and small amount. The Endocrine Culture

National Scientific Recommendations on Vitamin-D define insufficiency as 25 ng/mL of serum 26-OH-D, shortfall as 20–28 ng/mL, besides sufficient as >31 ng/mL.

The incidence of HCV infections in the Lahore province is roughly 23%, which would be worrying when contrasted to other areas of the country. As a result, preventative measures should be adopted in order to eradicate the current disease. Because vitamin D insufficiency is closely linked to HCV contagion, this effort remained meant to provide information about the current state of vitamin D in HIV - infected cases by evaluating incidence of 26 (OH) D3 among recent subgroups of Treated patients.

2. METHODOLOGY

Seventy-eight (n=78) participants (by means of an experiential technique) have been registered from June 25th, 2020 to February 25th, 2018 to learn about the relationship of Vitamin-D status on Hepatitis C virus. Our current randomized cross-sectional research was performed out at Lahore's Services Hospital. Blood samples were taken. All subjects provided consent forms. The sample was divided into two treatment populations: renumerated cirrhotic cases (n=24) to hepatitis C RNA-PCR positivity nonetheless no ascites, and fluid resuscitation cirrhotic sick people (n=28) through hepatitis C RNA-PCR positivity, ascites, and/or other cryptograms of illness of decompensation such as jaundice, hematemesis, and so forth. In furthermore, 25 seronegative for anti-HCV antibody normal subjects were included on the investigation. These individuals had no other documented illnesses and had not used Vitamin D supplements in the previous six months.

HCV RNA amplification and detection: QIAGEN GmbH's "QIA symphony SP" automated work station was used to extract HCV RNA. It operates on the ferromagnetic electron automatic RNA extraction approach.

Table 1.

Features	Control (n=26)	Compensated Cirrhosis (n=28)	Decompensated Cirrhosis (n=27)	P-value
Age (Mean, range)	37.08(22-54)	47.69(28-67)	51.68 (34-67)	< 0.001
Gender	9/14	10/12	10/19	0.729
Joint Pains (No/Yes)	6/17	2/20	9/20	0.166
Ascites (No/Yes)	22/1	22/0	29/0	< 0.002
Jaundice (No/Yes)	24/1	23/2	7/26	< 0.002
Hematemesis (No/Yes)	24/1	24/1	13/19	< 0.001
Blood Transfusion (No/Yes)	19/4	14/8	15/14	0.068
HCV RNA PCR QL (No/Yes)	23/0	0/22	0/29	< 0.002
Serum 25(OH)D ng/mL	31.4134	27.8506	21.6485	< 0.002

2.1 25-OH-Vitamin-D Measurement

The blood attentions of 25-OH-Vitamin-D in cases also normal remained determined that use the Virto's ECi immunodiagnostic instrument and an improved chemiluminescence technique. Sample, decoupling buffer, also conjugate buffer remained applied to a Virto's microwell decorated through just the monoclonal antibody in that order. To interact only with antibody, detached Vitamin-D contended through conjugate Vitamin-D. Unbound items were rinsed away after an incubation. The emission spectrum was again monitored after the addition of an increased chemiluminescence substrate. The data is updated monthly in nanograms per milliliter (ng/mL). Categorical data were reported in terms of percentages and frequencies, whilst quantitative variables generally provided in terms of mean SD.

For categorical variables, 3 tests in addition Kruskal- Walli's examinations remained rummage-sale to evaluate the differences among groups. Observed data without standard or asymmetrical distributions were evaluated through income of Student's t test or Mann-Whitney test. GraphPad Prism 8 was utilized to create figures.

3. RESULTS

As per the empirical technique, 78 people were adopted for this research. Subjects investigated divided into two categories based on clinical findings and anti-HCV titers: healthy controls (n=24) and hepatitis C cases (n=52). As indicated in Table 1, the HCV sample further

divided into 2 subcategories: compensation cirrhosis (n=23) besides decompensated cirrhosis (n=28). The individuals ranged in age from 21 to 66 years. HCV cases appeared younger than healthy controls (P0.001). The average age of the controls, HCV cases, and subcategories equaled 37.088.126 years.

The proportion of both control subjects having confirmed diagnosis (49.6913.446) and fluid resuscitation fibrosis (50.698.410) comprised females (61.3 percent and 61.9 percent respectively). Nonetheless, on the basis of sex, the differential was non-significant (p=0.728). Because the general population struggles from Vitamin-D deficiency, no significant changes in joint discomfort observed found (p=0.167). There used to be a substantial alteration in jaundice, hematemesis, overall blood transfusion activity (P 0.069) for biochemical markers (Table 1).

The 28 (OH) Vitamin-D level in HIV - infected cases found substantially lower than in healthy participants (Table 1; Fig.1). In HCV treatment groupings, there was no notable change in Vitamin-D levels between control and cHCV participants (P= 0.118) significantly (p versus dHCV (P= 0.002).

Whenever Vitamin-D distributions evaluated characterized using the Endocrine Society Professional Standards and guidance, the majority of the study group was determined to be deficient in Vitamin-D [6]. Vitamin-D deficit and inadequacy observed related among healthy controls, all HCV cases, and subgroups (P0.0002 for all group and P=0.02 control vs. HIV - infected cases) (Table-II; Fig.2).

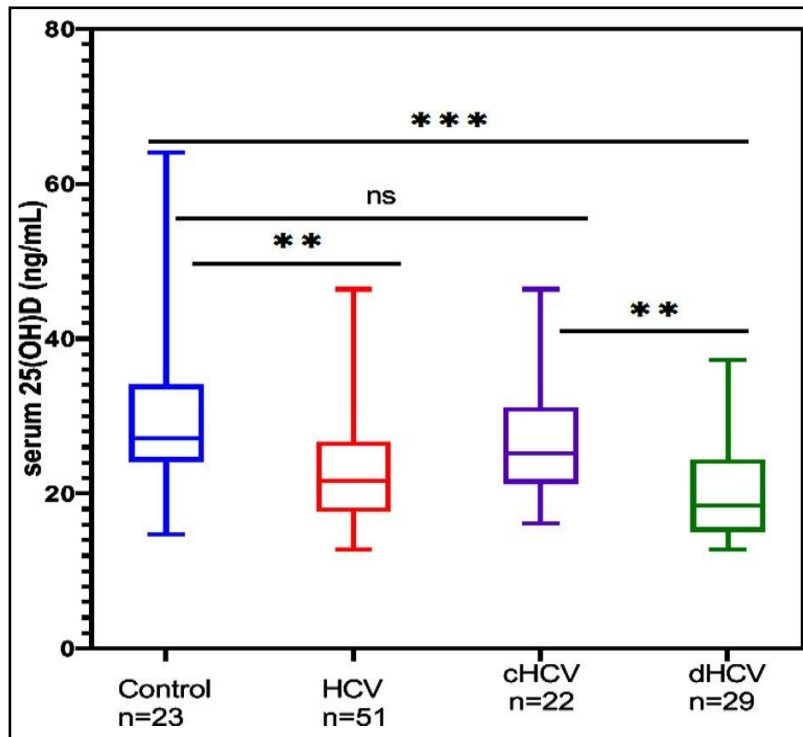


Fig. 1.

Table 2.

Vitamin-D stratum	Control (n=25)	HCV (n= 54)	Remunerated Cirrhosis (n= 28)	Decompensated Cirrhosis (n =25)
Scarce	3 (9.8%)	18 (38.4%)	4 (14.7%)	17(37.1%)
Insufficient	12(18.9%)	23 (38.3%)	13 (14.1%)	9(28.7%)
Normal	11 (23.1%)	13 (25.7%)	8(32.9%)	6(18.3%)

$P = <0.0002$

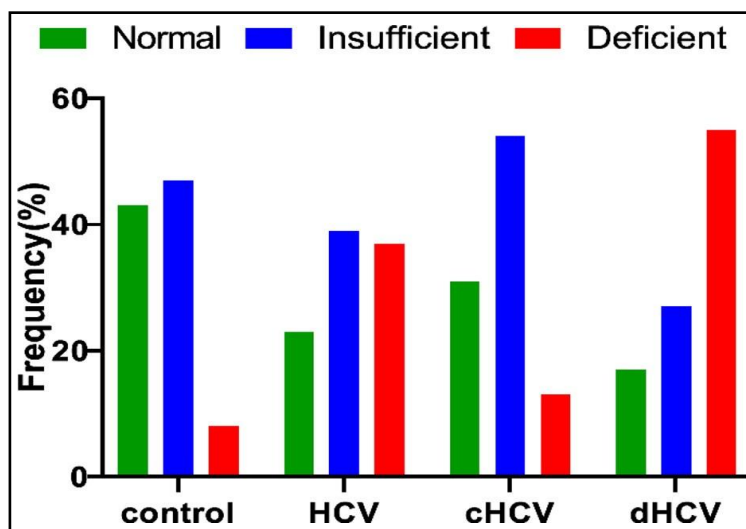


Fig. 2.

$P = 0.031$

4. DISCUSSION

Possession with the Hepatitis C virus is very serious public health concern that contributes significantly to the worldwide burden of liver problems. HCV infection is a primary cause of liver cirrhosis, which is the fourth leading cause of mortality in many parts of the globe. These have been found that those with chronic liver illness are more susceptible to acquire vitamin D insufficiency. Vitamin D's role is not limited to their conventional action in bone homeostasis in addition calcium control; the vitamin likewise remained engaged in the physiopathology of many illnesses as an immune modulator in both adaptive immunity immunological mechanisms [7]. Vitamin-D insufficiency has been observed to be prevalent in chronic liver disorders at a rate of around 93 percent.

As a result, the current research remained designed to evaluate status of 25 (OH) D in HCV cases divided into 2 sets based on liver tumor growth: compensatory liver cirrhosis and metabolic acidosis liver cirrhosis.

According to this research, 78.7 percent of HIV - infected patients had suboptimal levels of Vitamin-D (38.4 percent shortfall and 39.2 percent inadequate level, accordingly) as compared with healthy controls persons [8]. Those results significantly support the conclusions of previous reports on Vitamin-D insufficiency in individuals with liver cancer. Kumar et al. discovered that 86 percent of cirrhotic individuals had insufficient Vitamin-D levels. Nevertheless, contrary to these results, prior studies have shown that 91 percent of participants had low levels of Vitamin-D. (32 percent deficit in and 58 percent insufficient). This one was due to the fact that the intensity in addition prognosis of HCV create malfunction of internal Vitamin-D synthesis as the consequence of diminished 7-dehydrocholesterol synthesis. Another factor that may contribute to low blood Vitamin-D levels is individuals' restricted exposure to sunshine [9].

Vitamin-D levels have been shown to have an inverse association with more liver cirrhosis, with 56.3 percent of respiratory failure cirrhosis people undergoing from Vitamin-D deficit contrasted to 14.7 percent of compensated cirrhosis people undergoing from Vitamin-D deficiency (exposed in Fig. 2). Those data imply that blood Vitamin-D levels have a crucial role in the diagnostic workup of HCV infection. Vitamin-

D, being pluripotent in nature, remains engaged in immune modulation also anti-swelling properties. As a result, it decreases HCV core antigen and viral particle production. A lack of vitamin D could also result in increased viral load, inflammation, and liver damage. Vidor and colleagues, on other hand, found that vitamin D deficiency remained not linked to liver cirrhosis illnesses and consequences [10].

In HCV patients, we found a 37.4 percent deficiency of 25 (OH) D. In addition, Laredo et al. discovered 42 percent of inadequate levels in their research sample. Adorer as well as colleagues discovered a 58 percent deficiency in serum Vitamin-D. support our findings that vitamin D deficiency is common in individuals having Hepatitis C virus contagion. Though, Vitamin-D deficiency remained similarly detected in regular healthy persons. Riaz and his team conducted a larger scale research to establish the incidence of Vitamin-D insufficiency in Pakistan, that revealed that mainstream of inhabitants is vitamin-D deficient. And that is why both healthy people and HCV-infected patients experienced joint discomfort. A variety of variables, particularly ethnic disparities, BMI, and dietary patterns, may contribute to variability in inadequate Vitamin-D levels.

5. CONCLUSION

We found that HCV individuals had lower levels of Vitamin-D content than healthy controls; 78.6 percent of HIV - infected patients had suboptimal heights of Vitamin-D. The harshness of Vitamin-D insufficiency remained closely related to the intensity of the condition. The majority of individuals with Vitamin-D insufficiency was higher in the respiratory failure sample than in the supplemented cirrhotic class. In addition to HCV positive individuals, 26 (OH) D deficiency was reported in the healthy control group. As a result, it is advised that Vitamin-D estimate be performed in conjunction with the HCV screening test. Vitamin D supplementation and lifestyle changes can enhance overall nutritional Status therefore slow the course of liver cancer in HIV - infected patients. Having said that, more large-scale, multicenter, and long-term investigations are required to confirm those results.

CONSENT

It is not applicable.

ETHICAL APPROVAL

It is not applicable.

COMPETING INTERESTS

Authors have declared that no competing interests exist.

REFERENCES

1. World Health Organization. Rolling updates on coronavirus disease (COVID-19); 2020. [Cited 2020 Jul 31] Available: <https://www.who.int/emergencies/diseases/novel-coronavirus-2019/events-as-they-happen>.
2. World Health Organization. Timeline of WHO's response to COVID-19; 2020. [Cited 2020 Jul 31] Available: <https://www.who.int/news-room/detail/29-06-2020-covidtimeline>
3. COVID-19 Dashboard by the center for systems science and engineering (CSSE) at Johns Hopkins university; 2020. [Cited 2020 Jul 31] Available: <https://gisanddata.maps.arcgis.com/apps/opsdashboard/index.html#/bda7594740fd40299423467b48e9ecf6>.
4. World Health Organization. WHO coronavirus disease (COVID-19) dashboard; 2020. [Cited 2020 Jul 31] Available: <https://covid19.who.int/>
5. Daneshkhah A, Agrawal V, Eshein A, Subramanian H, Roy HK, Backman V. The possible role of vitamin D in suppressing cytokine storm and associated mortality in COVID-19 patients. MedRxiv. 2020;04.08.20058578. Available: <https://doi.org/10.1101/2020.04.08.20058578>.
6. Zhao X, Zhang B, Li P, Ma C, Gu J, Hou P, Guo Z, Wu H, Bai Y. Incidence, clinical characteristics and prognostic factor of patients with COVID-19: a systematic review and meta-analysis. MedRxiv. 2020.03.17.20037572. Available: <https://doi.org/10.1101/2020.03.17.20037572>.
7. Undela K, Gudi SK. Assumptions for disparities in case-fatality rates of coronavirus disease (COVID-19) across the globe. Eur Rev Med Pharmacol Sci. 2020;24(9):5180–2. Available: https://doi.org/10.26355/eurev_202005_21215
8. Bikle D. Vitamin D: Production, metabolism, and mechanisms of action. In: Feingold KR, Anawalt B, Boyce A, Chrousos G, Dungan K, Grossman A, Hershman JM, Kaltsas G, Koch C, Kopp P, Korbonits M, McLachlan R, Morley JE, New M, Perreault L, Purnell J, Rebar R, Singer F, Trencle DL, Vinik A, Wilson DP, editors. Endotext (Internet). South Dartmouth (MA): MDTText.com, Inc. 2000;76. Available: <https://www.ncbi.nlm.nih.gov/books/NBK278935/>
9. Christakos S, Dhawan P, Verstuyf A, Verlinden L, Carmeliet G. Vitamin D: Metabolism, molecular mechanism of action and pleiotropic effects. Physiol Rev. 2016;96(1):365–408. Available: <https://doi.org/10.1152/physrev.00014.2015>.
10. Colotta F, Jansson B, Bonelli F. Modulation of inflammatory and immune responses by vitamin D. J Autoimmun. 2019;85:78–97. Available: <https://doi.org/10.1016/j.jaut.2017.07.007>

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