Case-control study of serum vitamin D concentrations in hospitalised patients with COVID-19 and hospitalised controls suffering with respiratory tract infections of differing aetiology

Zakaria Ali Ibrahim Elmi,1 Sameer Sighakoli,1 John Tetteh,2 Nazanin Zand3

ABSTRACT
This study explored the prevalence of low serum vitamin D in patients admitted with acute respiratory tract infections (ARTIs) such as COVID-19. This study investigated whether patients with COVID-19 had lower serum vitamin D compared with patients with ARTIs of other aetiology. A case–control study was performed with cases of COVID-19 and controls of non-COVID-19 ARTIs. Patients were enrolled from a single general medical ward in a secondary care hospital between 15 April 2020 and 15 May 2020. Exclusion criteria were an oxygen requirement of >8 L/min. Data collected included serum 25-hydroxyvitamin D concentration, venous plasma glucose concentration and haemoglobin A1c. Outcomes measured were length of hospital stay, deaths, the need for high dependency and intensive care unit involvement. A total of 60 patients of five ethnic groups were enrolled, 85% (n=46) were of White-British ethnicity. The data analysis is based on these 46 patients of which 24 were non-COVID-19 patients with ARTI and 22 were patients with COVID-19. Overall, 80% of the study population had a serum vitamin D concentration below 50 nmol/L with median concentrations of 30 nmol/L and 35 nmol/L for patients with COVID-19 and non-COVID-19 ARTIs respectively. A Mann-Whitney sign-ranked test with respect to serum vitamin D concentration found no statistically significant difference between cases and controls, p=0.09. There was no significant difference in the length of stay, body mass index and rates of various comorbidities such as diabetes mellitus (DM), hypertension and lung disease in both study groups. However, DM was found to be associated with lower serum vitamin D concentrations. The results of this study support published literature showing an association between low serum vitamin D and ARTIs including COVID-19. However, this study did not identify patients with COVID-19 to have a statistically significant lower serum vitamin D concentration than non-COVID-19 patients with ARTI.

INTRODUCTION
SARS-CoV-2, a novel member of the coronavirus family, is the cause of the COVID-19. SARS-CoV-2 represents one of seven coronaviruses known to infect humans, and the third virulent strain causing severe pneumonia and acute respiratory distress syndrome (ARDS).1–3 Infections with SARS-CoV-2 have produced a wide spectrum of phenotypes, ranging from asymptomatic individuals to mild symptoms such as fever, dry cough, myalgia, diarrhoea and anosmia, to more severe disease manifestations including hypoxaemia, pneumonia, ARDS, multiorgan failure and death.4 ARDS is a common cause of mortality and morbidity in patients with COVID-19. A cytokine storm driven by the virus has been implicated. Patients with the infection have shown the upregulation of inflammatory cytokines such as IL-6.5 Similarly, overactivation of proinflammatory Th17 cells and cytotoxic CD8 T cells have been observed.6 The subsequent astounding release of proinflammatory
cytokines has the capability to induce systemic consequences such as shock and multiorgan failure.

In a similar manner to SARS-COV-1, SARS-CoV-2 is known to enter host cells via the cell surface receptor ACE 2 and reduce its expression in lung tissue.\textsuperscript{7,8} Experimental evidence from mice studies suggests that the manifestation of ARDS in relation to SARS COV-2 is partly due to a shift in the equilibrium away from ACE-2 and angiotensin 1–7 towards a predominance of ACE and angiotensin 2 resulting in more severe acute lung injury.\textsuperscript{9–11}

Vitamin D\textsubscript{3} (1,25(OH)\textsubscript{2}D)\textsubscript{3} is a fat-soluble secosteroid which typically participates in calcium homeostasis. Multiple studies have demonstrated an extraskeletal role of vitamin D in its interaction with the immune system and the renin-angiotensin-aldosterone system (RAAS).

Vitamin D receptor (VDR)-negative mice demonstrate elevated renin and angiotensin II production as well as more severe pulmonary oedema compared with wild-type mice following lipopolysaccharide aspiration.\textsuperscript{12,13} A multitude of adaptive and innate immune cells possess VDR. The exact immunoregulatory effects of vitamin D are often disease specific. The effects of vitamin D on cells of the innate immune system are largely stimulatory, resulting in enhanced phagocytoses and an amplified production of antimicrobial peptides.\textsuperscript{14} On the contrary, the effects of vitamin D on the adaptive immune system are characteristically inhibitory.\textsuperscript{15–18}

It is proposed then that low serum vitamin D concentrations may exacerbate COVID-19, first by worsening the already displaced equilibrium of ACE and ACE-2, and second, through the disinhibition of the adaptive immune system.

The risk factors associated with low serum vitamin D concentration are many. A cross-sectional survey of 1828 middle aged Caucasian adults identified lower serum vitamin D concentrations in relation to obesity/underweight, female gender, reduced physical activity and reduced sun exposure.\textsuperscript{19} A meta-analysis of 23 studies also found that obesity was associated with a 35% increased prevalence of vitamin D deficiency when compared with normal body weight individuals.\textsuperscript{20} The biological mechanisms underpinning the associations between body mass index (BMI) and vitamin D deficiency are yet to be understood. It has been suggested that obesity may result in decreased serum concentrations of vitamin D secondary to decreased sun exposure and increased fat sequestration.\textsuperscript{21,22}

A systematic review and meta-analysis of 17 observational studies consisting of 2756 patients identified an association between vitamin D deficiency and higher mortality, higher rates of hospital admission and prolonged patient stay in adults with COVID-19.\textsuperscript{23}

Another meta-analysis of 26 observational studies consisting of 8176 patients with COVID-19 found that vitamin D deficiency did not increase the risk of developing COVID-19; however, severe cases of COVID-19 were associated with vitamin D deficiency.\textsuperscript{24} A limitation to both of these meta-analyses included the inability to independently assess the impacts of confounding variables such as gender, ethnicity, BMI, diabetes and advanced age, all of which are known risk factors for vitamin D deficiency and severe COVID-19 outcomes.\textsuperscript{25,26}

Prior to COVID-19, vitamin D deficiency has been studied with particular interest in the context of acute respiratory tract infections (ARTIs). A 2017 individual participant data meta-analysis including 10933 participants in 25 randomised control trials found an overall protective effect of vitamin D supplementation against ARTI, the greatest benefit being seen in those with severe vitamin D deficiency.\textsuperscript{25}

These studies in combination with the biological plausibility reiterate the important immunomodulatory role of vitamin D in relation to ARTIs including COVID-19.

**AIM**

The aim of this study was to evaluate the vitamin D status of inpatients with COVID-19 and inpatients with non-COVID-19 ARTIs.

**METHOD**

**Research design**

Case–control study.

**Participants**

The demographic information for the participants included in the study are shown in **Table 1**. Participants included a total of 60 inpatients from a single general medical ward at Medway Maritime Hospital, Gillingham, Kent, United Kingdom between 15 April 2020 and 15 May 2020.

<table>
<thead>
<tr>
<th>Table 1: Participant demographics</th>
<th>COVID-19 +ve</th>
<th>COVID-19 –ve</th>
</tr>
</thead>
<tbody>
<tr>
<td>Total number of participants</td>
<td>30</td>
<td>30</td>
</tr>
<tr>
<td>Mean age (SD)</td>
<td>71 (SD 12.98)</td>
<td>69 (SD 12.51)</td>
</tr>
<tr>
<td>Number of men</td>
<td>18 (60%)</td>
<td>13 (43.3%)</td>
</tr>
<tr>
<td>Number of women</td>
<td>12 (40%)</td>
<td>17 (56.7%)</td>
</tr>
<tr>
<td>Number of White British patients</td>
<td>26 (87%)</td>
<td>26 (87%)</td>
</tr>
<tr>
<td>Number of Black British patients</td>
<td>0</td>
<td>1</td>
</tr>
<tr>
<td>Number of Black African patients</td>
<td>2</td>
<td>0</td>
</tr>
<tr>
<td>Number of Asian Indian patients</td>
<td>1</td>
<td>0</td>
</tr>
<tr>
<td>Number of patients for whom ethnicity was unrecorded</td>
<td>1</td>
<td>3</td>
</tr>
</tbody>
</table>
Exclusion criteria for admission to this particular ward included an oxygen requirement of over 8 L/min.

Procedure
Blood sampling and determination of serum vitamin D concentration
At admission participants had three blood samples taken: one serum separating tube, one fluoride oxalate tube and one potassium EDTA tube. These samples were processed for 25-hydroxyvitamin D, random venous plasma glucose and haemoglobin Alc (HbAlc).

25-hydroxyvitamin D was processed by UniCel DxI Immunoassay System using Beckman Access 25(OH) Vitamin D UniCel DxI Reagent Pack. Serum vitamin D concentration reference ranges used in this study are the National Institute for Health and Care Excellence (NICE) recommended ranges as follows: Less than 25 nmol/L is vitamin D deficiency, 25–50 nmol/L is vitamin D insufficiency and greater than 50 nmol/L is vitamin D sufficiency.

Plasma glucose and HbA1C determination
Enzymatic UV testing was used for the quantitative determination of glucose in human plasma on a Beckman Coulter AU analyser (AU680) using Beckman reagents. The glucose method used was a hexokinase/glucose 6 phosphate dehydrogenase method. The quality of all glucose results was assured by the continued participation in the United Kingdom National External Quality Assessment Service (UK NEQAS). High-pressure liquid chromatography was performed for HbA1C measurement using TOSOH BioScience system.

COVID-19 status determination
On admission to hospital, nasopharyngeal and oropharyngeal swabs were processed for the detection of COVID-19 using real-time reverse transcription PCR amplification technology performed on VitaPCR Instrument.

Auxiliary patient information
Information was collected from general practice (GP) and hospital records regarding patient’s admission date, discharge date, comorbidities, BMI, medications and any intensive care/high dependency stay during the admission.

Data analysis
Data selection
Initially, 60 patients met the non-exclusion criteria. It was observed that 87% of the patients were of White British ethnicity. The rest of the ethnic groups consisted of Black British, Asian Indian, Black African and those for whom ethnicity was unrecorded. Due to the sparse and statistically unrepresentative numbers, these groups were excluded from further analysis. The sample subjected to statistical analysis included only the White British discharged patients.

Principal component analysis:
The whole data set of 60 patients and seven variables consisting of random plasma glucose concentration, HbAlc, age, weight, height, length of hospital stay and serum vitamin D concentration were initially examined with Principal Component Analysis (PCA). PCA is a multivariate data analysis technique which enables the reduction of high dimensional data into a set of uncorrelated principal components that provide further insights into relationships between variables. Recent applications of PCA in COVID-19 studies have been published. A projection of serum vitamin D concentration (nmol/L), length of hospital stay (days) and BMI (kg/m²) onto the principal components graphically provided insights regarding the pattern distribution of patients relative to their COVID-19 status. Further information is provided in online supplemental material.

Statistical analysis
With respect to the 46 discharged patients of White British ethnicity, normality tests by Kolmogorov-Smirnov (p=0.000), Lilliefors (p=0.023) and Anderson-Darling (p=0.007) showed that the data was not normally distributed. The Mann-Whitney sign-ranked test was used for ascertaining p values.

Graphical analysis
Graphical profiles were generated with respect to COVID-19 status; serum vitamin D concentration; length of stay; BMI; and comorbidities such as diabetes mellitus (DM), hypertension (HTN) and lung disease (LD).

PCA and statistical analysis were performed in Matlab. Tabulations and visualisations were performed in Microsoft Excel and Tableau software.

RESULTS
Comorbidities
Table 2 shows that the comorbidities among both cohorts were of similar frequency. The total number of comorbidities was 37 in the COVID-19 +ve cohort and 41 in the COVID-19 –ve cohort.

Table 2 Frequency of comorbidities in the patient cohorts
<table>
<thead>
<tr>
<th>Comorbidity</th>
<th>COVID-19 +ve (n=22)</th>
<th>COVID-19 –ve (n=24)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Ischaemic heart disease</td>
<td>5</td>
<td>6</td>
</tr>
<tr>
<td>Atrial fibrillation</td>
<td>4</td>
<td>3</td>
</tr>
<tr>
<td>Heart failure</td>
<td>3</td>
<td>1</td>
</tr>
<tr>
<td>Chronic obstructive pulmonary disease</td>
<td>2</td>
<td>2</td>
</tr>
<tr>
<td>Diabetes mellitus</td>
<td>8</td>
<td>9</td>
</tr>
<tr>
<td>Chronic kidney disease</td>
<td>3</td>
<td>5</td>
</tr>
<tr>
<td>Hypertension</td>
<td>11</td>
<td>12</td>
</tr>
<tr>
<td>Asthma</td>
<td>1</td>
<td>3</td>
</tr>
<tr>
<td>Total number of comorbidities</td>
<td>37</td>
<td>41</td>
</tr>
</tbody>
</table>
of comorbidities exceeds the total number of patients highlighting that some patients had more than one comorbidity.

**Vitamin D analysis**

Table 3 highlights the degree of vitamin D insufficiency and deficiency seen in the entire hospitalised patient cohort. Irrespective of COVID-19 status, 80% (N=37) of the entire cohort had low serum vitamin D concentrations (<50 nmol/L) with 39% (N=18) having vitamin D deficiency.

The median vitamin D concentration for both study groups was within the insufficient range. A Mann-Whitney test revealed no statistically significant difference between the length of stay and serum vitamin D in both groups (p value 0.86 and 0.09, respectively).

The median serum vitamin D concentration for the patients who required high dependency unit or intensive treatment unit were paradoxically higher than those who did not. However, these numbers are too few to draw any statistically significant conclusion.

The histogram in figure 1 shows the left skew with respect to serum vitamin D concentration in both study groups.

**Length of stay analysis**

It is observed in figure 2 that the lowest p value of 0.122 was seen when comparing patients discharged with a length of stay shorter than 14 days to those with a length of stay longer than 14 days.

Figure 3 shows that no patient with a vitamin D serum concentration of greater than 50 nmol/L stayed longer than 14 days.

**Subgroup analysis with regards to BMI and comorbidity**

Figure 4 shows that the median BMI of COVID-19-positive patients of 25.3 was not significantly different from the median BMI of 27.8 seen in the COVID-19-negative patients. It can be noted that eight underweight patients (BMI<18.5) were seen in the COVID-19-positive group while only one was seen in the COVID-19-negative group, unfortunately such a small number does not lend itself to statistical comparison. No association was seen between

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**Table 3** COVID-19 positive and negative cohorts with regards to serum vitamin D concentration and length of inpatient stay

<table>
<thead>
<tr>
<th></th>
<th>COVID-19 +ve (n=22)</th>
<th>COVID-19 –ve (n=24)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Number of patients with vitamin D sufficiency (&gt;50 nmol/L)</td>
<td>2 (9%)</td>
<td>7 (29%)</td>
</tr>
<tr>
<td>Number of patients with vitamin D insufficiency (25–50 nmol/L)</td>
<td>9 (41%)</td>
<td>10 (42%)</td>
</tr>
<tr>
<td>Number of patients with vitamin D deficiency (&lt;25 nmol/L)</td>
<td>11 (50%)</td>
<td>7 (29%)</td>
</tr>
<tr>
<td>Median vitamin D (nmol/L)</td>
<td>30</td>
<td>35</td>
</tr>
<tr>
<td>Number of patients that required HDU/ITU</td>
<td>4</td>
<td>2</td>
</tr>
<tr>
<td>Median serum vitamin D concentration for HDU/ITU escalations (nmol/L)</td>
<td>32.5 (n=4)</td>
<td>51 (n=2)</td>
</tr>
<tr>
<td>Median length of stay (days)</td>
<td>6</td>
<td>7</td>
</tr>
</tbody>
</table>

HDU, high dependency unit; ITU, intensive treatment unit.
Figure 3  Serum vitamin D concentration with respect to length of stay irrespective of COVID-19 status. Horizontal lines highlight serum vitamin D concentrations of 50nmol/L and 40nmol/L.

BMI and length of stay, BMI and serum vitamin D concentration, BMI and COVID-19 status.

Figure 5 depicts the significant prevalence (37%) of DM in the study sample. There was no association between DM and length of stay or COVID-19 status. However, an association is noticed with regards to DM and serum vitamin D concentration irrespective of COVID-19 status. The majority of the patients with DM (88%) had serum vitamin D concentrations of less than 50nmol/L.

Figure 6 shows that hypertension was seen in 50% of the study sample. No obvious association was seen between hypertension and serum vitamin D concentration, hypertension and length of stay, hypertension and COVID-19 status.

Figure 7 shows lung disease (chronic obstructive pulmonary disease (COPD) and asthma) to have been present in 15% of the study sample. No obvious association was seen between lung disease and serum vitamin D concentration, lung disease and length of stay, lung disease and COVID-19 status.

DISCUSSION

The findings of this study demonstrate the significant prevalence of vitamin D insufficiency and deficiency in hospitalised patients with both COVID-19 and non-COVID-19 ARTIs. Eighty per cent of the study sample had a serum vitamin D concentration <50nmol/L. These results support previous studies demonstrating the presence of low serum vitamin D during ARTIs.25

Such prevalence would advocate for proactive screening of vitamin D, especially in those admitted to hospital for respiratory tract infections. It has been shown that improving overall vitamin D status could help reduce the economic burdens of several diseases.33

We did not observe a statistically significant difference in the serum vitamin D concentrations of patients with COVID-19 compared with non-COVID-19 patients with ARTI, both study groups had low median serum vitamin D concentrations of 30 nmol/L and 35 nmol/L, respectively.

The use of a control group consisting of a non-hospitalised healthy cohort may have produced a starker contrast in serum vitamin D concentration.

In this study, we found no statistically significant association between length of stay and serum vitamin D concentration. However, we did note that no patient with a normal serum vitamin D concentration had a length of...
CONCLUSION

This study was able to demonstrate the significant prevalence of vitamin D insufficiency and deficiency in hospitalised patients with ARTIs including COVID-19. This study did not demonstrate patients with COVID-19 to have a statistically significant difference in serum vitamin D compared with non-COVID-19 patients with ARTIs. Both study groups demonstrated near similar insufficiencies and deficiencies in vitamin D. Further studies comparing such groups could provide useful insights into the relationship of serum vitamin D concentration and ARTIs of various aetiologies including COVID-19.

Current literature on COVID-19 and vitamin D is suggestive of an association, a meta-analysis consisting of 39 studies of various designs and sample sizes, along with a meta-analysis of 10 case–control studies have shown vitamin D deficiency to be associated with increased risk of COVID-19 infection.36 37

Further work is required to ascertain whether there is a genuine causation underpinning this relationship.

Figure 7  Projection of lung disease (LD) on COVID-19 status relative to serum vitamin D concentrations and length of stay.

A large comparative analysis performed in the USA of 906,849 hospitalisations estimated the attributable risk of DM at 20.5%, obesity at 30.2% and hypertension at 26.2%.30 In this study, the substantial prevalence of hypertension and diabetes mellitus within the COVID-19-positive group was apparent at 50% and 37%, respectively.

The limitations of this study include its nature as an ethnically homogenous small sample. This limitation was driven by external factors such as sample collection window and rates of patient turnover. The study was also confined to moderately unwell cases of hospitalised patients requiring less than 8 L/min of oxygen further limiting its size.
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ORCID iD
Zakaria Ali Ibrahim Elmi http://orcid.org/0000-0003-0545-1989

REFERENCES