TABLE 1 (Continued)

Countries	Latitude degrees	Total cases (N)	Total deaths, N (%)	Deaths/1 Million population, N
Nigeria	10	174	2 (1.1)	0.02
Costa Rica	10	375	2 (0.5)	0.4
Ghana	8	195	5 (2.6)	0.2
Panama	8	1317	32 (2.4)	11
Venezuela	8	144	3 (2.0)	0.2
Ivory Coast	8	190	1 (0.5)	0.1
Sri Lanka	7	148	3 (2.0)	0.2
Cameroon	5	255	6 (2.3)	0.3
Malaysia	5	3116	50 (1.6)	2
Brunei	4	133	1 (0.7)	2
Colombia	3	1065	17 (1.7)	0.7
Singapore	1	1000	4 (0.4)	1
Ecuador	-1	2758	98 (3.5)	10
Indonesia	-2	1790	170 (9.5)	0.7
DRC	-4	123	11 (8.9)	0.2
Peru	-7	1323	47 (3.6)	3
Mayotte	-13	116	1 (0.9)	7
Bolivia	-17	123	7 (5.7)	0.9
Mauritius	-20	161	7 (4.3)	6
Australia	-25	5137	25 (0.5)	1
Southern Hemisphere				
South Africa	-29	1380	5 (0.4)	0.2
Chile	-31	3031	16 (0.5)	2
Argentina	-34	1133	33 (2.9)	1
Uruguay	-34	350	2 (0.6)	1
New Zealand	-41	797	1 (0.1)	0.1

Note: Data extracted from https://www.worldometers.info/coronaviru s/ (Accessed April 2, 2020).

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Letter: Covid-19, and vitamin D. Authors' reply

EDITORS,

We read with great interest the letter from Drs Panarese and Shahini, regarding our review article. We are glad that our review has helped Italian colleagues in this pandemic and grateful for their comments.

Interestingly, they found that northerly latitude is associated with increased mortality rate and hospitalisation rate for COVID-19 world-wide.¹ One possible explanation was the vitamin D deficiency due to low ultraviolet exposure in Northern countries and now probably compounded by "shut-down" measures, as mentioned by Drs Panarese and Shahini. It has also been suggested that weather conditions of

low temperature and relative humidity might allow the virus to survive longer outside the body than under warmer conditions.

A recent review that also supported the possibility of vitamin D reducing the risk of COVID-19 infections and deaths documented the various relevant actions of vitamin D.² These include maintenance of cell junctions, strengthening cellular immunity by reducing the cyto-kine storm with impacts on tumour necrosis factor α and interferon γ ,² and modulating adaptive immunity through suppressing T helper cell type 1 (Th1) responses and promoting induction of T regulatory cells.³ Vitamin D supplementation increases the CD4⁺ T cell count in

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HIV infection,⁴ and one of the main manifestations of severe SARS-CoV-2 infection was lymphopenia. Vitamin D deficiency can therefore be implicated in ARDS, and heart failure as well as sepsis,² and these can all be manifestations of critically ill COVID-19 patients.

It is also important that vitamin D is involved in two other critical regulatory systems. Thus hypovitaminosis D promotes the renin-angiotensin system (RAS), chronic activation of which may lead to chronic cardiovascular disease (CVD) and decreased lung function.⁵ Patients with these comorbidities account for a higher proportion of critically ill cases in COVID-19. Recently Hanff et al⁶ speculated that CVD or RAS blockade drugs might augment ACE2 levels, increasing available substrate for SARS-CoV-2 infection. SARS-CoV-2 infection is thought to downregulate ACE2 function, leading to toxic Angiotensin II overaccumulation which in turn may contribute to ARDS or fulminant myocarditis. Another prominent feature of severe COVID-19 is coagulopathy. A higher level of D-dimer was found in ICU patients than non-ICU patients,⁷ indicating a predominantly pro-thrombotic DIC. It was confirmed in pathology that microvascular thrombosis was found in lung tissues of COVID-19.8 Vitamin D deficiency has also been reported to correlate with increased risk for thrombosis, and vitamin D controls the expression of several genes relevant to cellular proliferation, differentiation, apoptosis, and angiogenesis.⁹

Therefore, together with Drs Panarese and Shahini, we agree that vitamin D deficiency may well be associated with an increased risk of severity in COVID-19. Further study of the impact of vitamin D levels on outcome in hospitalised patients is urgently needed. Meanwhile, it seems highly plausible that appropriate supplementation of vitamin D, as already recommended for populations with high prevalence of vitamin D deficiency, may reduce the risk of severe COVID-19 outcomes.

The authors' declarations of personal and financial interests are unchanged from those in the original article. 10

LINKED CONTENT

This article is linked to Sands et al papers. To view these articles, visit https://doi.org/10.1111/apt.15555 and https://doi.org/10.1111/apt.15716.



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Letter: corticosteroid use alongside tofacitinib in OCTAVE Open

Editors,

We read with great interest the study by Sands *et al.*¹ The authors found that flexible dosing with tofacitinib 5 mg b.d. and tofacitinib 10 mg b.d. can be incorporated into long-term

disease management strategies for patients with ulcerative colitis (UC). This study is important and interesting; however, one weakness of this study is the lack of information about whether glucocorticoid withdrawal was considered as a key secondary