


## LETTER

Thrombotic and immunological  
phenomena following COVID-19  
vaccination: experience from a case seriesChathurika Lakmini Dandeniya <sup>1</sup>, Theekshana Kavinda Premathilake<sup>2</sup>

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Vaccination has undoubtedly reduced COVID-19-related mortality globally.<sup>1</sup> However, being a new vaccine with limited clinical trial experience, postmarketing data are important to establish safety and side effect profile. Thrombotic and autoimmune phenomena have been reported globally following vaccination.<sup>2</sup> A multi-centre study in Sri Lanka failed to identify any significant adverse events following AZD1222 (ChAdOx1) first dose in Sri Lanka.<sup>3</sup> However, we present our experience with a case series identifying serious thrombotic and autoimmune complications following different types of COVID-19 vaccination in Sri Lanka.

We collected data from patients who presented to Teaching Hospital Peradeniya with thrombotic and/or immunological phenomena following COVID-19 vaccination from September 2021 to March 2022. Being a tertiary care hospital affiliated to a national university, the referred patients were from across the country. A causal relationship was assumed in patients in whom the symptoms appeared within 6 weeks of vaccination, without similar symptoms preceding the vaccination. Active COVID-19 infection was excluded in all patients with thrombotic complications and in all patients requiring hospital admission due to other symptoms. In patients presenting to outpatient rheumatology services, five had tested negative for COVID-19 infection around the time of symptom onset while the others did not display symptoms of infection around this time prompting testing according to national policy.

Of a total of 24 patients with their age ranging from 19 to 78 years, 18 were females. All had received at least one of the five types of COVID-19 vaccine used in

Sri Lanka at the time (BNT162b2, mRNA-1273, BBIBP-CorV, Gam-COVID-Vac and AZD1222 (ChAdOx1)) while four had received cross-vaccination. Eight patients (33.3%) mentioned stepwise progression of symptoms with booster doses. One patient died while the remaining 23 patients recovered with varying degrees of disability. Details of patients are presented in [table 1](#). Further details of all clinical cases can be accessed in online supplemental table 1.

The first patient with digital thrombosis was managed with apixaban and intravenous prostacyclin. However, the thrombosis progressed and later she developed thromboembolic pulmonary hypertension despite being on apixaban. Occult malignancy was considered but excluded. She succumbed to severe right heart failure 6 months later. Subsequent patients with digital thrombosis were tried on bosentan which was effective with rapid halt of progression of ischaemia and pain relief. All patients with inflammatory arthritis needed disease-modifying antirheumatic drugs due to failure of inducing sustained remission with steroids and non-steroidal anti-inflammatory drugs. All patients with immunological phenomena needed varying regimens of immunosuppression. Only two achieved complete remission with a short course of steroids while all others needed longer courses of immunosuppression.

Despite being rare, COVID-19 vaccination can be associated with unusual autoimmune and thrombotic phenomena which may intensify with repeated vaccination. For thrombotic complications, prostacyclin is of dubious efficacy while bosentan is effective. The majority needed sustained strong immunosuppression customised to the individual patient.

**Table 1** The observed clinical phenomena and outcomes

Clinical syndrome	Patients (n)	Treatment	Outcome
Distal digital thrombosis	3	Prostacyclin in one case. Bosentan in other cases. Oral prednisolone in all patients with 3 months of anticoagulation.	Prostacyclin was ineffective for halting progress of ischaemia while bosentan was very effective. The one on prostacyclin later developed thromboembolic pulmonary hypertension despite apixaban and died from complications.
Cerebral vasculitis: imbalance and cognitive deficit	2	Intravenous methylprednisolone in both. Intravenous immunoglobulin induction with azathioprine maintenance in one patient and intravenous cyclophosphamide induction with MMF maintenance in the other.	Full recovery to baseline with no new lesions on follow-up MRI.
De novo inflammatory arthritis: seropositive rheumatoid arthritis (n=2), peripheral spondyloarthritis (n=6)	8	All patients relapsed on stopping initial prednisolone or NSAIDs with persistent symptoms beyond 3 months, needing DMARDs.	All achieved remission.
Worsening of previously diagnosed autoimmune rheumatological conditions: SLE (n=2), rheumatoid arthritis (n=2), peripheral spondyloarthritis (n=1)	5	4 out of 5 were in long-term remission with good drug compliance. One patient with SLE had active disease with intermittent flares but in remission for 7 months preceding vaccination. All patients needed escalation of immunosuppression.	All achieved remission.
Cutaneous vasculitis with inflammatory oligoarthritis (positive ANA)	1	Leflunomide (intolerance of methotrexate).	Achieved remission.
Isolated cutaneous vasculitis	1	Prolonged course of tapering prednisolone 0.5 mg/kg/day.	Achieved remission.
Severe autoimmune haemolytic anaemia with positive ANA	1	Intravenous methylprednisolone effective but relapse on oral prednisolone. Remission induction with rituximab. Maintenance with hydroxychloroquine.	Achieved remission.
Unusual cognitive symptoms with positive ANA: dissociation and paraesthesia	1	Moderate-dose prednisolone tapered over 3 weeks.	Achieved remission without sequela.
RS3PO-like picture	1	Prednisolone, methotrexate, leflunomide.	Unable to completely stop steroids. Remission with 2 DMARDs and prednisolone 5 mg every other day.
Complex picture of digital ischaemia, cutaneous vasculitis and mucosal ulceration	1	Prednisolone, warfarin, bosentan.	CMV serology negative. Digital ischaemia stopped progression rapidly with bosentan. Remission with cessation of all medications after 1 year of treatment. Lost 4 digits.

ANA, antinuclear antibody; CMV, cytomegalovirus; DMARDs, disease-modifying antirheumatic drugs; MMF, mycophenolate mofetil; NSAIDs, non-steroidal anti-inflammatory drugs; RS3PO, remitting seronegative symmetrical synovitis with pitting oedema; SLE, systemic lupus erythematosus.

The findings of this observational study have to be interpreted within the limitations of the inability to completely rule out asymptomatic COVID-19 infection as a potential cause in all the cases (done in some cases only), inability to rule out preceding undiagnosed autoimmune disease or occult malignancy, consideration of the contribution from comorbidities and the possibility of natural flares of baseline rheumatological disease.

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**Ethics approval** This study involves human participants but this is a case series instead of a planned and executed study. Data were collected from patients with their written consent as and when they presented with these unusual and rare manifestations. Therefore, ethics approval was not sought prior to starting data collection and it was not practically possible. Participants gave informed consent to participate in the study before taking part.

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