Dear Editor,

The coronavirus disease 2019 (COVID-19), classified as a pandemic by the World Health Organization (WHO), has affected nearly all countries and regions around the world. Evidence to date suggests that patients with concomitant medical conditions are at higher risk for severe COVID-19 (1). However, data regarding the comorbidities in children with COVID-19 is relatively rare (2). We report a case of suspected COVID-19 infection in a 6-year-old boy with Henoch–Schonlein purpura (HSP) complicated by diffuse alveolar hemorrhage.

On April 1, 2020, the patient was brought to the emergency department of Bahrami Children’s Hospital, Tehran, Iran, because of a palpable purpuric maculopapular rash involving his left ankle and foot and signs of arthritis in the left ankle and knee. The rest of the physical examination was unremarkable. The child was admitted for further evaluation. Laboratory tests showed leukocytosis, increased ESR and CRP and mild hematuria and proteinuria. The patient was clinically diagnosed with HSP, was given supportive treatment and ibuprofen. Arthralgia and edema gradually subsided and the patient was discharged on day 3 with a prescription of ibuprofen.

Two days later, the patient was readmitted with fever, tachypnea, dyspnea, 02 saturation of 94% on room air, bilateral fine rales on lung auscultation and abdominal pain associated with melena. Patient’s hemoglobin was decreased and urine analysis showed hematuria and proteinuria. Abdominal ultrasound demonstrated mural thickening of distal ileum and decreased peristalsis. Chest X-ray (CXR) showed bilateral, peripheral ground glass opacities of the lungs. COVID-19 rapid test was negative; however, a clinical diagnosis of COVID-19 was made and after PICU admission treatment started with methyl prednisolone, cloxacillin, azithromycin and hydroxychloroquine. As patient’s condition worsened and his hemoglobin level dropped to 6.1 gr/dL, packed red cells were given. COVID-19 rapid test repeat revealed negative results. The patient was suspected to have developed pulmonary hemorrhage associated with HSP. Bronchoscopy and bronchoalveolar lavage (BAL) showed diffuse alveolar hemorrhage. As BAL specimens tested negative for SARS-CoV-2, the patient received intravenous cyclophosphamide. However, treatment was not successful and the patient died of respiratory failure.

COVID-19 is a major concern to rheumatologists as rheumatologic patients may potentially be at an increased risk of infection and death due to the underlying immune dysfunction and treatment with immunosuppressive agents (3). However, data regarding rheumatologic diseases as a risk factor for increased incidence or severity of COVID-19 in children is very scarce (4).

Few reports have described adults and children presenting with maculopapular, purpuric and acro-ischemic skin lesions who were subsequently diagnosed with COVID-19 (5, 6). Moreover, vasculitis-like lesions due to endothelial damage are reported in severe forms of COVID-19 (7). At first presentation, our patient had the typical palpable purpuric skin lesions of HSP and his symptoms were successfully managed with ibuprofen. However, he was readmitted with pulmonary manifestations, which led to a clinical diagnosis of COVID-19, and exacerbation of HSP-related symptoms.

Despite negative COVID-19 rapid test, our primary treatment strategy was to manage both HSP and COVID-19 manifestations, i.e. methyl prednisolone, antibiotics and hydroxychloroquine. With worsening of respiratory manifestations, unexplained decline of hemoglobin level led us to perform bronchoscopy and BAL, which confirmed diffuse alveolar hemorrhage, a rare complication of HSP which manifests with tachypnea, dyspnea, sudden drop of hemoglobin and diffuse pulmonary infiltrates and is associated with poor prognosis (8).
During hospitalization, our patient was tested three times for COVID-19, one taken from the BAL specimen. It is believed that severe respiratory distress met in critically-ill COVID-19 patients results from a cytokine storm syndrome, a condition which may also complicate rheumatologic diseases, including vasculitis. At the time of the epidemic, patients presenting with severe respiratory manifestations and underlying vasculitis should be suspected of COVID-19 despite a primary negative COVID-19 rapid test.

This report doesn’t draw any conclusions on the possible association between pediatric rheumatologic conditions and COVID-19. Obviously, all children suspected with COVID-19 should be managed according to the standard protocols but it is mandatory to investigate other differential diagnoses, particularly in patients with underlying conditions. In case of concurrent COVID-19 and a rheumatologic condition, it seems reasonable to choose a treatment plan to manage both conditions.

Clinical judgment and decision making about vasculitis is challenging in COVID-19 pandemic due to:

1) COVID-19 can mimic cutaneous, pulmonary and renal manifestations of vasculitis.
2) COVID-19 can trigger some vasculitic signs and symptoms.
3) Vasculitis treatment influenced by COVID-19 pandemic while sensitivity and specificity of RT-PCR and COVID-19 serology are unclear.

Footnotes

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