Dear Sir,

The role of real-time polymerase chain reaction (RT-PCR) is pivotal in the diagnosis of severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2). The COVID-19 pandemic started from Wuhan, China, in December 2019, and the disease is still evolving.\(^1-3\) False-negative RT-PCR results have been reported in human immunodeficiency virus (HIV) patients with COVID-19.\(^4\) However, data on the features of COVID-19 in these patients is scarce.\(^5\) We present a case of co-infection of HIV-1, herpes simplex virus-2 (HSV-2) and SARS-CoV-2 with false-negative RT-PCR in a critically ill patient.

The first case of COVID-19 in our country, Saudi Arabia, was confirmed on 2 March 2020, and more than 316,000 people had been infected and more than 3,900 patients had died at the time of writing.\(^6,7\) In May 2020, at the peak of the COVID-19 outbreak in Saudi Arabia, a previously healthy 40-year-old woman was admitted to the emergency department with four days of fever (38.5°C), persistent cough, myalgia and dyspnoea. Her peripheral oxygen saturation was 70% on room air, and she had haemodynamic instability. Her score on the Glasgow Coma Scale was 9 out of 15, with no other neurological pathology. She was intubated, mechanically ventilated and transferred to the intensive care unit (ICU) designated for COVID-19. Laboratory findings were normal, apart from lymphocytopenia (0.69 × 10\(^9\)/L; normal range [NR] 1.1–3.2 × 10\(^9\)/L), and increased levels of C-reactive protein (706 mg/L; NR 0–5 mg/L) and lactate dehydrogenase (990 units/L; NR 100–190 units/L). The rest of the biochemical report was within normal limits.

The patient underwent a full diagnostic work-up for viral, bacterial and systemic disorders. Cardiac enzymes, electrocardiogram and echocardiography results were normal. Chest computed tomography (CT) depicted bilateral ground-glass opacities and consolidations (Fig. 1a). She was tested for SARS-CoV-2 infection using RT-PCR assays (targeting for the RdRp gene, E gene and N gene of SARS-CoV-2), which were performed with nasopharyngeal swabs using the QuantiNova Probe RT-PCR kit (Qiagen, Valencia, CA, USA) in a Light-Cycler 480 real-time PCR system (Roche, Basel, Switzerland).\(^8,9\) Specifically, each 20 μL of reaction mixture contained 10 μL of 2 × QuantiNova probe RT-PCR master mix, 0.2 μL of QN Probe RT-Mix, 1.6 μL of each 10 μM forward and reverse primer, 0.4 μL of 10 μM probe, 1.2 μL of RNase-free water and 5 μL of TNA as the template. The thermal cycling condition was ten minutes at 45°C for reverse transcription, five minutes at 95°C for PCR initial activation, and 45 cycles of five seconds at 95°C and 30 seconds at 55°C. The assays were performed as previously described.\(^10\) However, RT-PCR results for SARS-CoV-2 were negative.

Emergency central nervous system (CNS) CT and angiography revealed diffuse white matter hypodensities and brain oedema (Fig. 1b & c). The differential diagnosis included stroke, encephalitis and posterior reversible encephalopathy syndrome, among others. Upon ICU admission (Day 1), lopinavir/ritonavir (400 mg/100 mg, two tablets of 200 mg/50 mg twice daily), acyclovir (10–15 mg/kg intravenously every eight hours), ceftriaxone (1 g intravenously twice daily), dexamethasone (10 mg intravenously once daily), therapeutic low-molecular-weight heparin (adjusted for the patient’s body weight and renal function) and ICU supportive care were administered by the treating ICU team.\(^11\) The patient also underwent the acute respiratory distress syndrome net protocol and received prone positioning ventilation, maintaining a partial arterial pressure of oxygen to fractional inspired concentration of oxygen ratio of 160. Cerebrospinal fluid (CSF) analysis revealed slightly elevated opening pressure of 22 cm CSF (NR 10–20 cm) and 205 white blood cells/μL (85% lymphocytes), with normal protein and glucose levels. PCR in the serum detected HIV-1 (plasma viral copies 3,000/μL; CD4 and CD8 T cells 350/μL and 684/μL, respectively). In the CSF, HSV-2-DNA was found by means of the HSV-1/2 PCR kit (Qiagen). Interestingly, another RT-PCR test for SARS-CoV-2 performed on nasopharyngeal specimens (24 hours apart) was persistently negative.

Fig. 1 (a) CT image of the chest shows bilateral diffuse peripheral ground-glass opacities and consolidations with mainly lower lobe distribution. (b) Brain CT and (c) CT angiography images show diffuse white matter hypodensities in the frontal-parietal region, extending to the corpus callosum, basal ganglia and brain stem (mainly on the left side).

Co-infection of human Immunodeficiency virus, herpes simplex virus-2 and SARS-CoV-2 in a patient with false-negative real-time polymerase chain reaction results

On Day 3, the patient developed septic shock and multisystem organ failure. Antibiotics were upgraded to meropenem (2 g intravenously every eight hours) and vancomycin (1 g intravenously every 12 hours). Vasopressors and fluid resuscitation were initiated. The patient developed acute kidney injury with associated acidosis and hyperkalaemia, and underwent continuous renal replacement therapy as per Kidney Disease Improving Global Outcomes 2019 guidelines. On Day 4 after ICU admission, the third RT-PCR test performed for COVID-19 was positive (31 copies for the E gene and 29 copies for the RdRp gene assays per reaction). Rescue therapies were considered; however, the patient developed ventricular tachycardia and cardiac arrest. No electrolyte disturbances were observed. Despite strenuous resuscitation efforts, the patient succumbed to the disease. Blood cultures for common bacterial and fungal infections were negative. CSF Cryptococcus and blood Toxoplasma gondii antigen serology tests were also negative. Cultures derived from endotracheal tube aspirates were negative for common bacteria, and microscopy examination, which was used for detection of Pneumocystis species, was also negative. In addition, tests performed for systemic disorders (i.e. autoimmune disorders, including antiphospholipid antibodies) were negative.

This brief case report, albeit with limitations, carries important messages. Life-threatening COVID-19 can present with acute respiratory distress syndrome, sepsis, multisystem organ failure, neurological manifestations and thromboembolic phenomena. Notably, devastating CNS pathology, ranging from meningoencephalitis to stroke and acute disseminated encephalomyelitis, has been reported in patients with severe COVID-19. In our patient, contrast brain CT did not depict any ring enhancement; however, the possibility of other diagnoses, such as toxoplasmosis or CNS lymphoma related to HIV, cannot be excluded. Unfortunately, magnetic resonance imaging was not performed as the patient deteriorated rapidly and could not be transported safely outside of the ICU. Also, the possibility of stroke remains a differential diagnosis and the detection of HSV-2 in the CSF cannot be ignored. This is a rare (<2%) cause of HSV encephalitis and has a bad prognosis, although it is rare for HSV-2 encephalitis to present with brain oedema and mass effect. We are uncertain how our patient was infected with SARS-CoV-2 and HIV, as her past medical history and list of conducts were unavailable. Hence, the exposure risk could not be properly evaluated, which is a major limitation of this case report, preventing its generalisability. Nevertheless, our patient had a relatively high HIV viral load with preserved CD4 count and CD4/CD8 ratio. Therefore, we speculate that our patient could have had acute HIV infection or HIV seroconversion illness, which is associated with CNS pathology, although its link to COVID-19 is not yet elucidated.

In conclusion, co-infection of SARS-CoV-2 with other viruses may result in unpredictable dysregulations of the immune system response that are not fully understood. The diagnostic and therapeutic challenges are obvious. Further research is required to investigate co-infections of SARS-CoV-2 with other viruses, as well as the diagnostic performance of RT-PCR in such cases.
References