Diaphragmatic paralysis in COVID-19: a rare cause of postacute sequelae of COVID-19 dyspnoea

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SUMMARY
We describe a 56-year-old female patient hospitalised with COVID-19 in April 2020 who had persistent respiratory symptoms after radiographic and microbiologic recovery. X-ray of the chest demonstrated an elevated right hemidiaphragm while fluoroscopy confirmed unilateral diaphragmatic paralysis. Symptoms resolved gradually, concurrent with restoration of right hemidiaphragm function. Thus, we describe a rare cause of postacute sequelae of COVID-19 dyspnoea.

BACKGROUND
COVID-19, caused by SARS-CoV-2, is a multisystem viral illness predominantly affecting the respiratory system. SARS-CoV-2 also targets the nervous system and patients can present with anosmia, dizziness, seizures, or, rarely, acute cerebrovascular disease.1 Neurologic symptoms are a consequence of direct viral effects on the central nervous system (CNS) or peripheral nervous system (PNS) and collateral damage from the inflammatory response. Case reports have described mononeuritis multiplex, Guillain-Barre syndrome and bilateral diaphragmatic paralysis associated with acute COVID-19; however, it is unclear whether PNS lesions affect patients after recovery from acute illness.2–4 Postacute sequelae of COVID-19 (PASC) affect up to two out of three of patients, up to 60% of which describe persistent dyspnoea more than 60 days after infection.5 Here, we describe a case of COVID-19-related unilateral diaphragmatic paralysis causing PASC dyspnoea.

CASE PRESENTATION
A 56-year-old woman with a body mass index of 32 and medical history of obstructive sleep apnea (OSA), hypertension, hyperlipidaemia and breast cancer presented in April 2020 with progressive dyspnoea, cough, headache, fever and chills after being diagnosed with COVID-19 by nasal swab 6 days earlier. Her admission vital signs were significant for a temperature of 101.1°F, heart rate of 104 beats per minute and oxygen saturation of 95% on room air. X-ray of the chest (XRC) demonstrated bilateral peripheral interstitial infiltrates (figure 1A).

She was admitted to the medical ward and administered supplemental oxygen. On day 5 of admission, she developed progressive hypoxaemia requiring admission to the intensive care unit for close monitoring. She was started on empiric ceftriaxone and azithromycin. On day 7 of admission, an XRC demonstrated worsening multifocal airspace disease consistent with COVID-19 acute respiratory distress syndrome (ARDS) and an elevated right hemidiaphragm (figure 1B). She was administered vancomycin and cefepime, furosemide, and was enrolled in a trial of tocilizumab for COVID-19 ARDS. The patient attempted to prone, but only tolerated lying on her side, without any preference for one side or the other. Her neck was not immobilised. She never received treatment with non-invasive ventilation or intubation and mechanical ventilation. Central venous access was deferred, and she never received steroids, remdesivir or vasopressors. She refused treatment for her OSA throughout her hospitalisation. Her hypoxaemia gradually improved, she was weaned off supplemental oxygen by day 15, and was discharged home on day 20. After discharge, the patient had persistent dyspnoea, dry cough and ambulatory hypoxaemia.

INVESTIGATIONS
The patient was referred to a pulmonologist who obtained a repeat XRC demonstrating resolution of pulmonary infiltrates and persistently elevated right hemidiaphragm (figure 1C). At the time of her first clinic visit, pulmonary function tests were obtained that demonstrated a restrictive pattern (forced expiratory volume in 1 second (FEV1) 1.88 L (76% predicted), forced vital capacity (FVC) 2.18 L (69% predicted), FEV1/FVC 87%, total lung capacity 3.64 L (76% predicted), residual volume 1.56 L (85% predicted)). CT of the chest showed normal pulmonary parenchyma and a persistently elevated right hemidiaphragm. Chest fluoroscopy demonstrated paradoxical right hemidiaphragm movement consistent with unilateral diaphragmatic paralysis. Reduced access to ambulatory testing during the pandemic prevented further investigation of diaphragm function (eg, electromyography, diaphragmatic ultrasound, seated/supine spirometry). A repeat XRC 11 months after discharge showed resolution of her right hemidiaphragm elevation (figure 1D).

OUTCOME AND FOLLOW-UP
The patient’s symptoms resolved over the following year with expectant management. She is now able to perform her own activities of daily living again and walk longer distances without getting short of breath.

DISCUSSION
PASC affects up to two out of three of COVID-19 survivors and commonly manifests with respiratory

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surgery, damage to the cervical nerve root, neuralgic amyotrophy and can be due to stretching/cooling during cardiothoracic surgery. Two possible processes could explain this phenomenon: direct neuroinvasion or an inflammatory process involving phrenic neuropathy. Our case suggests that SARS-CoV-2 could be an additional viral aetiology of SARS-CoV-2 infection directly affecting the PNS from a non-neuronal reservoir is also possible. A study of diaphragm muscle obtained at autopsy demonstrated SARS-CoV-2 infection of diaphragm myofibers, suggesting this could be a potential source for phrenic nerve retrograde invasion. A second explanation is that SARS-CoV-2 could have caused mononeuritis multiplex of the phrenic nerve. Two potential mechanisms could induce mononeuritis multiplex. First, a SARS-CoV-2-induced prothrombotic state or systemic vasculitis led to the formation of microthrombi within the vasa nervorum, inducing ischaemia and neuronal dysfunction. Second, release of proinflammatory cytokines during SARS-CoV-2 infection directly impaired neuronal function.

This case report adds important dimensions to our understanding of diaphragmatic paralysis in COVID-19. One prior report describes diaphragmatic paralysis in a patient infected with COVID-19 with a history of OSA who was treated with continuous positive airway pressure (CPAP) and mechanical ventilation. A second case report describes respiratory failure in an obese patient with COVID-19 attributable to diaphragmatic paralysis. Our patient is obese and has OSA, suggesting these may be risk factors for the development of diaphragmatic paralysis with COVID-19. A preprint describing a cohort of 25 patients with COVID-19 treated with mechanical ventilation identified a high burden of diaphragmatic abnormalities. Another study describes 1527 patients with COVID-19, 1.5% of which were diagnosed with diaphragmatic dysfunction on CT scan. These reports differ from ours as each patient was treated with CPAP or mechanical ventilation, both known risk factors for the development of diaphragmatic dysfunction, while our patient received no non-invasiveventilation or mechanical ventilation. Taken together, our report suggests that OSA and obesity may be risk factors for the development of diaphragmatic paralysis during COVID-19 infection.

Treatment for unilateral diaphragmatic paralysis is reserved for symptomatic patients and involves weight loss and/or nocturnal positive pressure ventilation. Patients with severe or bilateral diaphragmatic paralysis may require diaphragmatic fundoplication. Our patient’s unilateral paralysis resolved with expectant management keeping with its likely viral origin.

**Learning points**

- We report a case of postacute sequelae of COVID-19 (PASC) dyspnoea due to unilateral diaphragmatic paralysis.
- This presentation is likely due to direct effects of SARS-CoV-2 on phrenic nerve function.
- Peripheral neuropathies should be considered as a cause of PASC-related dyspnoea.
- Obesity and obstructive sleep apnea could be risk factors for development of diaphragm paralysis in the setting of COVID-19 infection.

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