## NEUROIMAGING HIGHLIGHT

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## Posterior Reversible Encephalopathy Syndrome in Acute COVID-19 Pneumonia

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A 66-year-old woman presented to the Emergency Department with shortness of breath and was found to be febrile and tachycardic. Chest CT showed bilateral ground-glass opacities, and she was admitted with a diagnosis of COVID-19 pneumonia. PCR testing confirmed that she was positive for the SARS-CoV-2 virus. Medical co-morbidities included asthma, type 2 diabetes, hypertension, and dyslipidemia. Shortly after admission, she was intubated and transferred to the ICU, where she remained for 40 days. There were no sustained episodes of hypertension. Bacterial superinfection led to Pseudomonas pneumonia, and her course was further complicated by subclavian deep vein thrombosis.

As she began to regain consciousness, she was confused and disoriented with headache, complaining that she could not see. Family members found that she did not recognize them when they entered the room, only when she could hear their voices. Two non-contrast CT scans of the brain were performed and were both interpreted as normal. When vision blurring persisted, MRI brain with diffusion-weighted imaging (DWI) was performed and showed extensive bilateral areas of T2/FLAIR hyperintensity (Figure 1), consistent with vasogenic edema and posterior reversible encephalopathy syndrome (PRES). Over the ensuing weeks, her vision recovered completely.

Acutely ill patients with COVID-19 are at risk for neurologic complications including stroke, viral meningitis/encephalitis, cerebral venous thrombosis, hypoxic encephalopathy, and demyelinating disease. PRES must be included in the differential diagnosis, and appropriate neuroimaging in the form of MRI with DWI performed. PRES is a result of an incompletely understood process of failed endothelial regulation resulting in cerebral vasogenic edema and was initially described in patients with immunosuppressive drugs, renal insufficiency, or severe hypertension. Occipital and parietal lobes are most frequently affected, and symptoms include headache, altered sensorium, seizure, and vision loss. Prompt treatment to reverse the causative stimulus results in recovery of function in most cases.

In COVID-19, the mechanism of PRES is most likely related to the viral-induced cytokine storm. One likely effector is tumor necrosis factor-α (TNFα), a major inflammatory cytokine released primarily by macrophages but also by many other cell types to incite multiple downstream signaling cascades and induce effects such as increased vascular permeability.<sup>2</sup> Other cytokines such as interleukin-1 likely play a role and upregulate cell adhesion molecules including intercellular adhesion molecule-1 (ICAM1) and vascular cell adhesion protein-1 (VCAM1).<sup>6,7</sup> An alternative hypothesis for the mechanism of PRES in these patients is dysregulation of the renin-angiotensin system (RAS) as SARS-CoV-2 spike protein binds to the angiotensin-converting enzyme-2 (ACE2) receptor to initiate cell entry.8 This leads to ACE2 downregulation and downstream effects on the RAS, and bradykinin-kallikrein pathway may cause endothelial dysfunction and increased vascular permeability. 9 Drugs used to treat COVID-19 including hydroxychloroquine and tocilizumab may also be associated with PRES, 10,11 though not a factor in this case.

PRES must be considered in COVID-19 patients with neurologic symptoms, particularly with severe bilateral vision loss. MRI of the brain with DWI is recommended in these cases to arrive at the correct diagnosis.

## DISCLOSURES

The authors have no conflicts of interest to declare.

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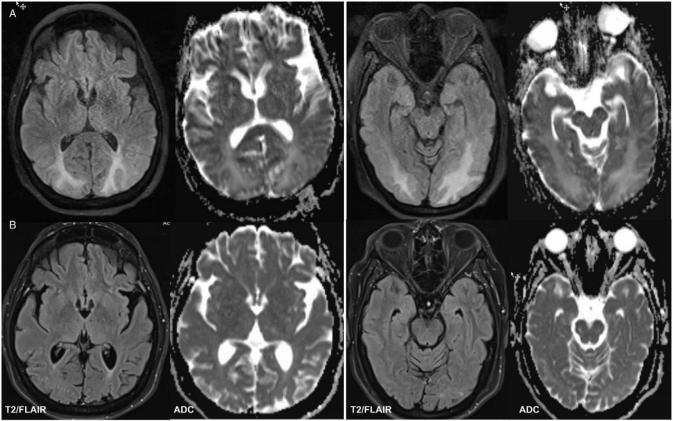


Figure 1: Posterior reversible encephalopathy syndrome in COVID-19. (A) Axial MRI images showing extensive bilateral regions of increased signal on T2/FLAIR sequences (left panels parieto-occipital, right panels occipital) and high signal on apparent diffusion coefficient sequences (ADC) consistent with extensive vasogenic edema. Most regions showed T2 shine through on diffusion-weighted imaging with sparse patches of restricted diffusion. (B) Follow-up MRI images showing resolution of edema.

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