The patient described in this report was the second child of healthy, non-consanguineous Korean parents. At 11 years of age, she was admitted to our clinic with fever, headache, and hemiparesis on the left side. On a neurologic examination, she showed facial palsy and decreased motor power (grade III) on the left side. Laboratory tests revealed elevated lactic acid (3.3 mmol/L). Brain magnetic resonance imaging (MRI) and angiography showed diffuse cortical and subcortical T2/fluid-attenuated inversion recovery (FLAIR) high signal intensity with gyral swelling in the right temporo-parieto-occipital lobe with steno-occlusive lesions (Supplementary Fig. 1). Mitochondrial DNA sequencing revealed m.3243A > G heteroplasmy of the mitochondrially encoded TRNA-Leu (UUA/G)1 (MT-TL1) gene, and her condition was diagnosed as mitochondrial encephalopathy, lactic acidosis and stroke-like episode (MELAS) syndrome (Supplementary Fig. 1). She presented with severe sensorineural hearing loss (SNHL) on pure-tone audiometry and a decreased full-scale intelligence quotient of 52 based on an intelligence test. Laboratory tests revealed elevated lactic acid (3.3 mmol/L). Brain magnetic resonance imaging (MRI) and angiography showed diffuse cortical and subcortical T2/fluid-attenuated inversion recovery (FLAIR) high signal intensity with gyral swelling in the right temporo-parieto-occipital lobe without steno-occlusive lesions (Supplementary Fig. 1). After administration of high-dose arginine therapy, she was discharged on admission day 6 with improved verbal responses. One week later, she was readmitted to our medical center because of dysarthria, poor verbal response, and general weakness. Repeated brain MRI showed increased extent of the previous lesions (Supplementary Fig. 2). During hospitalization, lip twitching was observed, and electroencephalography showed diffuse suppressed background activity with intermittent delta slow waves over the left frontal areas (Supplementary Fig. 2). After conservative therapy, including arginine and gabapentin, her dysarthria and lip twitching gradually disappeared. On hospitalization day 12, she returned to her usual condition...
and was discharged.

Here, we present a case of new stroke-like episodes after COVID-19 in a patient with MELAS syndrome. This case provides additional information regarding COVID-19 and could guide the evaluation and management of patients with mitochondrial disease during the current pandemic situation. Since the initial outbreak of severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2), previously referred to as “the novel coronavirus,” in December 2019, there is mounting evidence that COVID-19 causes not only respiratory illness but also a hypercoagulable state, leading to thrombotic complications including ischemic stroke [1]. In a recent review by the World Stroke Organization, comprising stroke experts from 18 countries, the risk of ischemic stroke in hospitalized patients with COVID-19 was approximately 5%, and the median duration from the first symptoms of COVID-19 to stroke was 10 days [2]. Suggested mechanisms of the development of ischemic stroke in COVID-19 include vasculitis, a hypercoagulable state, and cardiomyopathy [1]. Additionally, Saleh et al. [3] suggested that mitochondrial dysfunction could occur in COVID-19. Stroke-like episodes are a cardinal feature of MELAS syndrome, occurring in 84% to 99% of patients along with seizure and lactic acidemia [4]. As in our patient, the most common cause of MELAS syndrome is a missense mitochondrial DNA mutation (m.3243A > G) in the MT-TL1 gene, encoding mitochondrial transfer RNA for leucine; this mutation reduces mitochondrial protein synthesis and energy production [5]. Energy deficiency stimulates the proliferation of abnormal mitochondria in smooth muscles and small blood vessel endothelial cells, causing angiopathy, cellular dysfunction, and microvasculature perfusion disorders, leading to segmental vasodilation defects and subsequent vasogenic edema, which are thought to contribute to the occurrence of stroke-like episodes [6-7]. The drug, chickenpox infection, and fever have been reported as potential triggering factors of stroke-like episodes [8]. Our patient with MELAS syndrome was relatively stable, without stroke-like episodes, and was doing well, except for SNHL and mild intellectual disability. However, the stroke-like episodes recurred after COVID-19. It is difficult to identify the exact pathomechanism, although one possibility is that the COVID-19 infection collapsed the compensatory state of preexisting mitochondrial dysfunction, energy failure, neuronal cytopathy, and angiopathy, and then the inflammatory response of vascular endothelium and additional mitochondrial dysfunction led to the stroke-like episode in our patient. Although there are accumulating reports of acute cerebrovascular disease as a complication of COVID-19, the development of stroke during COVID-19 is more frequent in patients with vascular risk factors, including diabetes mellitus (odds ratio [OR], 5.56; 95% confidence interval [CI], 3.34 to 9.24), hypertension (OR, 7.35; 95% CI, 1.94 to 27.87) and coronary artery disease (OR, 3.12; 95% CI, 1.61 to 6.02) [9]. We need to raise awareness of mitochondrial encephalopathy as a risk

Fig. 1. Brain magnetic resonance imaging of first admission at 15 years of age.
factor for stroke in COVID-19 and provide appropriate guidelines to these patients.

**Supplementary materials**

Supplementary materials related to this article can be found online at https://doi.org/10.26815/acn.2021.00395.

**Conflicts of interest**

No potential conflict of interest relevant to this article was reported.

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Methodology: MJK. Project administration: WJK. Visualization: WJK and MJK. Writing-original draft: WJK. Writing-review & editing: MSY, MJK, and TSK

**References**