



# A case of hemichorea following administration of the Pfizer-BioNTech COVID-19 vaccine

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Dear Editor,

Hemichorea is a rare condition presenting involuntary hyperkinetic movement on the unilateral side and is primarily caused by structural damage in the basal ganglia or subthalamic nucleus. Some cases of hemichorea have been caused by cortical or thalamic lesions [1]. Though the structural damage is induced mainly by a vascular lesion, hemichorea can develop from other causes including infection, inflammation, and metabolic disease. As many people have received vaccines against coronavirus disease 2019 (COVID-19), various adverse effects (AE) have been reported. The Pfizer-BioNTech COVID-19 vaccine, a messenger RNA (mRNA) vector vaccine, is the only vaccine approved by the U.S. Food and Drug Administration (FDA) and has fewer AEs than vaccines with other platform technologies [2]. However, inflammatory diseases including myocarditis, pancreatitis, and lymphadenopathy and various symptoms have been reported after administration of the Pfizer-BioNTech vaccine [3]. We present a patient who developed hemichorea after administration of the Pfizer-BioNTech vaccine.

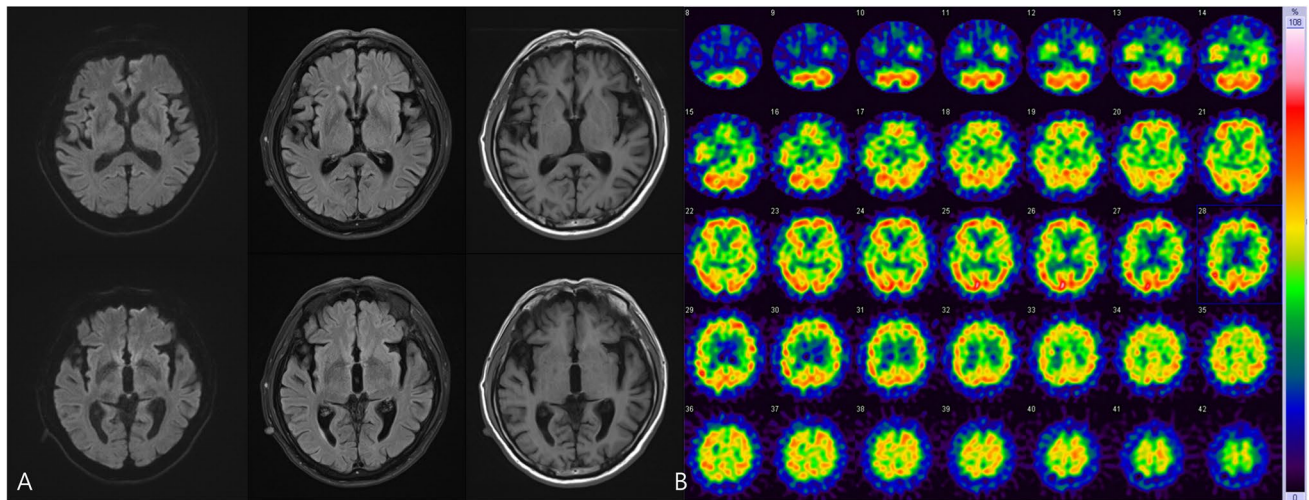
An 83-year-old male was referred for neurologic evaluation due to involuntary hyperkinetic movement of his right arm and leg for 1 month. It was slow, irregular, and purposeless movement and had occurred within 1 day after administration of the second dose of the Pfizer-BioNTech COVID-19 vaccine ([supplementary video](#)). He had developed no AE after the first dose of the vaccine at 3 weeks prior to the second. His cranial nerve functions, motor power, sensory and cerebellar functions were unremarkable. He denied other symptoms including fever, headache, and myalgia and had

neither DM nor history of stroke, although he did take some medicines for hypertension and benign prostate hyperplasia. Brain magnetic resonance imaging (MRI) performed one and half months later from the onset of the symptom showed normal findings except mild stenosis of the middle cerebral and basilar arteries (Fig. 1A). Brain SPECT demonstrated the perfusion pattern asymmetrically decreased in left thalamus (Fig. 1B). Laboratory tests for anti-streptolysin-O, serum copper, serum ceruloplasmin, vitamin B12, folate, anti-nuclear antibody, anti-phospholipid antibody, C3 and C4 complements, and RA factor were normal, as were HbA1C (5.6%), D-dimer (0.61 mg/L), and prothrombin time INR (1.04). Electroencephalography showed symmetric brain activity without abnormal activity. We started haloperidol 0.75 mg two times a day, and his symptom was mostly relieved at two weeks of follow up.

The Pfizer-BioNTech COVID-19 vaccine can impart immunity against severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) using an mRNA vector to deliver the genetic code of the antigen to a host. AEs of the Pfizer-BioNTech COVID-19 vaccine have been reported less frequently and at less severe levels than in vaccines with other technical platforms [2]. Most AEs with the Pfizer-BioNTech COVID-19 vaccine are simple local reactions and systemic reactions including fever, myalgia, and headache. Serious AEs, such as myocarditis, lymphadenopathy, and anaphylaxis, have been reported in rare cases. Neurological AEs that developed after administration of the Pfizer-BioNTech COVID-19 vaccine were tingling, numbness, limb weakness, vertigo, seizure, and loss of consciousness. Bell's palsy and Guillain–Barre syndrome were reported in some cases [3, 4]. Brain structural damage has been rarely reported after the Pfizer-BioNTech COVID-19 vaccine. Although further studies are in process to understand the mechanism of AE in the Pfizer-BioNTech COVID-19 vaccine, it is supposed that inflammation or infection in multiple organs including those of the nervous system could occur after the vaccination.

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**Fig. 1** Brain MRI results of the patient (A). Diffusion-weighted, fluid-attenuated inversion recovery, and T1-weighted axial images show normal findings in the basal ganglia, thalamus, and surrounding

brain regions. Brain SPECT (B) showed left thalamic hypoperfusion compared with right side

Headache and severe AEs have been reported with myocarditis and anaphylaxis.

The pathophysiology of hemichorea in this case is unclear. Acute onset of the hyperkinetic movement on the unilateral side supposes that it is caused by structural damage in the brain. Brain SPECT revealed contralateral thalamic hypoperfusion, which can be concerned with hemichorea. Although no structural damage was found on brain MRI, a small lesion can be concealed between slices 4-mm thick. In addition, metabolic disturbances induced by the Pfizer-BioNTech COVID-19 vaccine could affect the thalamus, which contains the regulatory pathway for movement. Thalamic dysfunction can disrupt important connections for the coordination of movement and develop aberrant pathway which can evoke abnormal involuntary movements. In ChAdOx1 adeno-virus vector vaccine, sudden increasing serum glucose level following vaccination has been reported in some cases [5]. In addition, other systemic illness may induce embolism, which can cause thalamic dysfunction or hypoperfusion.

To the best of our knowledge, this is a first report of hemichorea following administration of the Pfizer-BioNTech COVID-19 vaccine. Although brain MRI, electroencephalography, and laboratory test showed no causative abnormality in this report, functional disturbance was found in contralateral thalamus using SPECT. His symptom was a typical aspect of hemichorea and developed suddenly in temporal association with the

vaccination. Therefore, it is supposed that the Pfizer-BioNTech COVID-19 vaccine can be cause of hemichorea by an interruption in the unilateral thalamic function. Due to the possibility that other factors could affect the development of hemichorea in this case, we planned to evaluate for follow up. Further studies and more reports like ours are needed to expand the knowledge of hemichorea and COVID-19 vaccines.

**Supplementary Information** The online version contains supplementary material available at <https://doi.org/10.1007/s10072-021-05763-5>.

## Declarations

**Ethical approval** None.

**Conflict of interest** The authors declare no competing interests.

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