Bilateral Tonic Pupils and Guillain–Barre Syndrome in a 6-Year–Old Boy

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Bilateral tonic pupils can be a rare manifestation of Guillain-Barré syndrome (GBS). Here we report our experience of bilateral tonic pupils in a child with GBS, who diagnosed by pharmacological pupil test with diluted 0.125% pilocarpine and nerve conduction velocity (NCV) study.

A 6-year-old boy was transferred to our hospital due to altered mental status after respiratory resuscitation. One day before the transfer, he had gastrointestinal symptoms and weakness for dehydration. Few hours before the transfer, he abruptly started vomiting and developed difficulty in breathing. Upon arrival at the emergency room, he was intubated and mental status was stupor (Glasgow coma scale E2V2M5, temperature 37.7°C, blood pressure 110/65 mm Hg, respiration rate 25/min). Both pupils were dilated > 5 mm and light reflexes were hardly detected bilaterally. He was flaccid and deep tendon reflexes were decreased.

He was born at full term by uncomplicated delivery (birth weight 4,000 g). Before admission, he had been healthy and neurodevelopment was normal. Medical and family histories were unremarkable. He had not been vaccinated recently. The initial laboratory results for blood, stool, and cerebrospinal fluid (CSF) were unremarkable, except for a mild hyponatremia (Table 1). Chest radiographs revealed right upper lobe atelectasis and peribronchial infiltrations, indicating aspiration pneumonia. Brain diffusion magnetic resonance imaging (MRI) at the emergency room and regular brain and spinal MRI scans on the fifth day showed no brain or spinal cord injury. There was no enhancement or thickening of upper and lower cranial nerves on brain MRI. Electroencephalography revealed diffuse cerebral dysfunction on the first day. Due to possibility of encephalopathy, hypoxic injury, or severe pneumonia, intravenous (IV) antibiotics and IV acyclovir were started, and the respiratory function and intracranial pressure increase signs were carefully monitored. Additionally, considering the possibility of rapidly aggravating GBS, we initiated intravenous immunoglobulin (IVIG) treatment from the second day (400 mg/kg/day for 5 days).

Within a few days, the patient became alert and his respiratory functions improved. However, the generalized motor weakness and pupil dilatation persisted. Ptosis or anhidrosis was not noted. The weakness was more severe in the lower (grade 2) than in the upper extremities (grade 3). On the 5th day, he was extubated. The pupil dilatation slowly improved to 5 mm, but the light reflex was weak and present only in the

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root enhancement mechanism. Adjunctive spinal MRI may demonstrate spinal nerve tiganglioside antibodies have been related with the pathogenic GBS, but cannot be detected at the early phase. Serum or CSF an Albumino-cytologic dissociation in the CSF is a characteristic of diplopia, can also occur nomic or cranial nerve involvement, such as facial weakness and involvement and some are intubated at the peak of illness. Auto infections or vaccination, and paralysis presents as ascending pat in boys estimated between 0.4 and 1.3/year/100,000 children and is higher 2 months after discharge from the hospital. He was doing well in dai right eye. Ocular movements were normal; he could watch movies on his smartphone and track moving objects. We performed a pharmacological test on the pupils using diluted 0.125% pilocarpine drops. By dropping the pilocarpine for an hour, pupillary con striction was induced, which was not observed in normal condition (Fig. 1). This finding corresponded to bilateral tonic pupils. The patient’s motor functions gradually recovered from upper to lower limbs, and pupil size and reflexes normalized. Rehabilitation consultation was performed and physiotherapy was started. On the 16th day, he could walk without assistance. NCV study was performed on the 16th day, and the results showed F-wave absence, which is consistent with bilateral tonic pupils. GBS is one of the most common causes of acute flaccid paralysis in children. It is an acquired polyradiculoneuropathy characterized by ascending weakness and areflexia. The incidence is estimated between 0.4 and 1.3/year/100,000 children and is higher in boys [1]. The symptoms often develop following common infections or vaccination, and paralysis presents as ascending pattern. Nearly 20% of the patients experience respiratory muscle involvement and some are intubated at the peak of illness. Autonomic or cranial nerve involvement, such as facial weakness and diplopia, can also occur [1].

The diagnosis of GBS is mainly based on the clinical findings. Albumino-cytologic dissociation in the CSF is a characteristic of GBS, but cannot be detected at the early phase. Serum or CSF antiganglioside antibodies have been related with the pathogenic mechanism. Adjunctive spinal MRI may demonstrate spinal nerve root enhancement [1]. During the subacute period, electromyography or NCV may reveal pathological changes—F response absence, reduced conduction velocity by demyelination, or reduced compound action potential amplitude in the axonal forms [1]. In case of suspicion of GBS, children should be hospitalized until stabilization. IVIG should be started as soon as possible. After 2 weeks of a plateau period, most patients gradually recover with patterns inverse to those of the progression. However, younger age and rapid progression may be correlated with long-term complications [1]. Rare severe fulminant GBS manifested as locked-in syndromes reported in an infant with herpes simplex virus infection [2].

In the present case, the patient initially thought as flaccid paralysis with aspiration pneumonia of unknown etiology. During the treatment, his mentality recovered soon. His clinical feature immediately before respiratory failure suggested ascending paralysis and he recovered as descending pattern. The NCV test result at subacute stage and bilateral tonic pupil further supported the diagnosis of a form of GBS [3,4].
Bilateral tonic pupils are caused by abnormalities of the ciliary ganglion or a postganglionic parasympathetic branch of the ciliary nerve [3,4]. In adults, bilateral tonic pupils are frequently idiopathic. Generalized peripheral neuropathy, autonomic neuropathies, diabetes mellitus, paraneoplastic syndromes, and other generalized diseases can manifest bilateral tonic pupils [4]. In GBS, demyelination of the postganglionic ciliary nerve can be manifested as tonic pupils. It may occur without extraocular muscle paralysis [4]. However, this condition is rare in adults and extremely rare in children [2,3,5]. The youngest child reported was an 11-year-old boy [5]. Diagnosis is based on instillation of diluted pilocarpine (0.0625% to 0.125%); it causes a supersensitive pupillary constriction in GBS, which does not appear in normal conditions [3,4]. In the present letter, we report a very rare case of acute bilateral tonic pupils in a child with GBS. Postganglionic involvement of the parasympathetic nerves may be the underlying cause. The pharmacological pupil test with diluted 0.125% pilocarpine was helpful for the diagnosis.

Written informed consent was obtained from parent. This study was approved by the Institutional Review Board at Hanyang University Guri Hospital (2019-07-023-002).

Conflicts of interest

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

References