A Complex Cortical Malformation Caused by a Mutation in the Tubulin-Encoding TUBB3 Gene

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Tubulinopathy commonly refers to complex congenital and non-progressive brain malformations caused by mutations in the tubulin genes. Among tubulin-encoding genes, TUBB3 has rarely been reported as a cause of complex cortical malformations. Herein, we report a case of tubulinopathy in a 21-month-old boy who presented with delayed development. He could not walk on his own and was not able to speak more than five words. Physical examination revealed right esotropia and hypotonia of the lower extremities. MRI showed dysmorphic brainstem and dysmorphic and hypertrophic basal ganglia. The right thalamus was relatively smaller than the left one. The cerebellum showed disorganization of the cerebellar folia. DNA sequencing revealed a missense mutation of the TUBB3 gene.

Index terms Nervous System Malformations; Brain; Neuroimaging; Tubulin; Microtubules

INTRODUCTION

Microtubules, composed of tubulin are essential for neuronal migration and axon guidance in developing brain. Tubulinopathy is mutations in tubulin genes commonly cause complex congenital and non-progressive brain malformations. Among tubulin-encoding genes, TUBB3 gene has rarely been reported as cause of complex cortical malformation. Here, we report a case of 21-month-old boy with TUBB3 gene mutation presented with delayed development.
CASE REPORT

A 21-month-old boy was admitted to our rehabilitation medicine department with complaints of delayed development. He could stand with holding by himself but not walk by himself and he wasn’t able to speak more than five words. On physical exam, he had right esotropia, poor dynamic stability of proximal muscles during gait and muscle power weakness of lower extremities. He was the second child of healthy parents, and his older sister was healthy. He was born at 37 gestational weeks with a birth weight of 2670 g (5th percentile) and his perinatal history was uneventful. Bayley scales of infant development revealed 11 months old in cognitive

Fig. 1. Brain imaging of a 21-month-old boy with delayed development reveals complex cortical malformations.
A. The hypertrophic left basal ganglia shows fusion between the putamen and the caudate nucleus with an indiscernible anterior limb of both the internal capsules (arrows). The left thalamus is relatively larger than the right thalamus (dashed arrow). The lateral ventricles are asymmetric (arrowheads).
B. The hypertrophic right basal ganglia shows fusion between the putamen and the caudate nucleus with an indiscernible anterior limb of both the internal capsules (arrows). An iso- to slightly high signal intensity lesion at the right peritrigonal white matter indicates a thickened external sagittal striatum or heterotopia (dashed arrows).
C. Dysmorphic brainstem with atrophy of the right midbrain (arrow) and disorganization of the cerebellar folia (dashed arrows) are seen.
D. The right pons shows atrophic changes (arrow).
E. The right middle cerebellar peduncle shows atrophic changes (arrow).
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MRI showed dysmorphic and hypertrophic basal ganglia with fusion between the putamen and caudate nucleus with indiscernible anterior limb of both internal capsules (Fig. 1A, B). Asymmetry of both lateral ventricles (Fig. 1A) and both thalamus were seen (Fig. 1A). An iso to slightly high signal intensity lesion at the right peritrigonal white matter indicated thickened external sagittal striatum or heterotopia (Fig. 1B). A disorganization of cerebellar folia was seen (Fig. 1C). The brain stem was dysmorphic with atrophy of right sided brainstem (Fig. 1C, D) and right middle cerebellar peduncle (Fig. 1E). These findings suggested complex cortical malformation caused by tubulopathy.

The malformation of cortical development targeted sequencing revealed a missense mutation of TUBB3 gene, c.646G>A (p.Glu216Lys) (Table 1).

DISCUSSION

Tubulins compose microtubule, a filamentous structure, which is critical to the processes of mitosis, axon navigation, and neuron migration, all key factors in brain development (1). Thus mutations in tubulin genes can alter the normal function and structure of microtubules, leading to complex congenital and non-progressive disorders of brain development characterized by severe brain malformations (2). Disorders caused by mutations in the tubulin gene family were recognized as tubulopathy.

Among tubulin encoding genes, diseases caused by mutations in TUBB3 are rare, and only 15 TUBB3 mutations have been identified (3). TUBB3 encodes beta tubulin isotype 3, a neuron-specific component of microtubules. Heterozygous missense mutations in TUBB3 have been reported to be associated with decreased microtubule stability (4).

Clinical symptoms result from hypoplasia of the oculomotor nerve are esotropia, nystagmus and paralytic eye movement disorder, such as congenital fibrosis of extraocular muscle, CFEOM3 (5). Patients show gross motor delays associated with axial hypotonia, sensorimotor polyneuropathy and intellectual disability (4).

In radiologic finding, tubulopathy results in multiple malformations including microcephaly, lissencephaly, cerebellar hypoplasia, band heterotopia and other types of cortical dysgenesis due to impaired neuronal migration. Impaired axonal pathfinding lead anomalies of white matter pathways and anomalies of cranial nerves. Other findings are hypoplasia or absence of the corpus callosum, a small asymmetric brain stem, fused striatum due to absence of various parts of the internal capsule (most commonly the anterior limb). These are some characteristic features to be recognized on routine MR imaging (1).

In our case, the patient had symptoms indicated tubulopathy, such as right esotropia, hypotonia of lower extremities, intellectual disability and delayed development. MRI showed dysmorphic basal ganglia with fusion of the caudate and putamen and hypoplastic and dysmor-

| ACMG Classification | Gene | Accession | Nucleotide | Amino Acid | Zygosity | Inheritance |
|---------------------|------|-----------|------------|------------|----------|-------------|
| Likely pathogenic   | TUBB3| NM_001197181.1 | c.646G>A | p.Glu216Lys | Hetero | AD          |

ACMG = American College of Medical Genetics and Genomics, AD = autosomal dominant
phic brain stem. Additionally in our case, the cerebellum showed disorganization of cerebellar folia. In our knowledge this finding was not yet reported.

Based on our review, the MRI findings play essential role in diagnosis of tubulinopathy. Thus, in a patient with combination of delayed development, dysgyria, dysmorphic basal ganglia and brain stem, the tubulinopathy should be considered.

Author Contributions
Conceptualization, P.N.H.; supervision, P.N.H.; writing—original draft, L.Y.H.; and writing—review & editing, P.N.H.

Conflicts of Interest
The authors have no potential conflicts of interest to disclose.

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