Pontocerebellar Hypoplasia Type 1 with Ohtahara Syndrome: A Case Report

Abstract
Pontocerebellar Hypoplasia type 1 is a very rare form of infantile spinal atrophy which has been known to be associated with seizures, however, this is the first report of Early Infantile Epileptic Encephalopathy (Ohtahara syndrome) with tonic seizures, which is an important association to recognize.

Keywords: Pontocerebellar Hypoplasia type 1; Electroencephalography; MRI; Emergency; Spinal Cord

Introduction
We present a case of Pontocerebellar Hypoplasia type 1, a very rare form of infantile spinal atrophy, and report its association with tonic seizures and burst suppression pattern EEG.

Case Presentation
A 5 day old male newborn delivered at 40 weeks gestation by emergency C-section due to fetal distress with Apgar scores of 5 and 6, at 1 and 5 minutes respectively, was noted to be hypotonic at birth with joint contractures. He required resuscitation and endotracheal intubation due to apnea at birth. On day 4 of life he developed tonic seizures and was treated with the anticonvulsant, phenobarbital with good seizure control.

The baby was the born to a Yemeni consanguinous couple, the mother was G4P3, there was a family history of 2 prior pregnancies that each resulted in death at age 4 months each with a similar clinical picture of weakness and joint contractures, while 1 prior pregnancy resulted in a male with a clinical picture of epilepsy and joint contractures who is now 9 years old.

Testing
An MRI of the brain (Figure 1) showed features of poor gyration, as well as abnormal appearing white matter with a leukodystrophic appearance and profound pontocerebellar hypoplasia. An EEG was done which showed a burst suppression pattern (Figure 2); a diagnosis of Pontocerebellar hypoplasia type 1 (PCH type 1) was made and appropriate counseling provided to the parents. Genetic testing was not performed.

Discussion
Pontocerebellar hypoplasia type 1 with infantile spinal atrophy (OMIM 607596) is a genetically heterogeneous rare autosomal recessive disorder. It typically has dramatic atrophy of the pontocerebellar structures readily recognizable by MRI imaging, along with a progressive course reminiscent of spinal muscular atrophy [1]. The first report of this condition by Norman, after whom the condition is named, described cerebellar hypoplasia in Werding-Hoffman disease, and he hypothesized that the association may be linked to a single genetic abnormality [2]. This has proven to be the case indeed, as cases of pontocerebellar hypoplasia with infantile spinal atrophy have been linked to homozygous mutations in the vaccinia-related kinase 1 (VRK1) gene [3] as well as TSEN54 and RARS2 genes [1].

Unlike previously reported cases, this child additionally presented with tonic seizures and a burst suppression EEG pattern that define Ohtahara syndrome (Early Infantile Epileptic encephalopathy with burst-suppression) [4]. To our knowledge this is the first and only report of Ohtahara syndrome in PCH type 1. Although the prognosis is poor in this progressive neurodegenerative disorder, it is important to recognize this possible neurological condition since visual MRI diagnosis in the context of the classic presentation is virtually diagnostic, when the majority of the cases have no genetic locus identified. Moreover, early recognition and management of seizures may help improve the overall clinical outcome, although unlikely to have an impact on long term prognosis.
Figure 1: Sagittal (A), Coronal (B,C) and axial T2 MRI brain images shows profound ponto-cerebellar atrophy. Axial T1 (E) and T2 (F) show simplified gyral pattern and leukodystrophic white matter.

Figure 2: Double banana montage routine EEG showing burst suppression pattern. (compressed 30 seconds/per page).
Conclusion

To our knowledge this is the first reported case to show the association of Early Infantile Epileptic Encephalopathy (Ottahara Syndrome) with Pontocerebellar atrophy type 1. It is important to recognize PCH type 1 as a possible underlying etiology for Ottahara syndrome, also, the presence of Ottahara syndrome should help distinguish PCH type 1 from other types of spinal muscular atrophy.

References


