Diffuse Leptomeningeal Glioneuronal Tumor: Case Report and Review of Literature

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Abstract
In this study, we describe a rare case of disseminated CNS malignancy of childhood. A seven-year-old boy with headache, vomiting and gait disturbances was diagnosed with non-obstructive hydrocephalus and papilledema. Initial MRI revealed no significant findings, however subsequent studies revealed bilateral extradural and leptomeningeal enhancement. Biopsy was eventually obtained with pathology report confirming leptomeningeal oligodendrogliomatosis. We discuss the recent description in literature of this rarely reported diagnosis. This entity primarily affects pediatric patients. It has a unique clinicopathological identity, which can be easily mistaken for an inflammatory process. The diagnosis of this tumor can be complicated due to the rather nonspecific clinical presentation and findings on imaging.

Keywords
Oligodendrogliomatosis, Leptomeningial spread, CNS malignancy, Glioneural tumor

Introduction
Disseminated CNS malignancies of childhood are rare. Until recently only a few cases were described in the literature. In recent years, however, reports on pediatric cases with distinct pathological and molecular features led to the creation of a new tumor entity called diffuse leptomeningial glioneuronal tumor [1-5].

The clinical diagnosis of this tumor is sometimes complicated due to the rather nonspecific imaging findings [6]. On MRI these lesions usually present as diffuse thickening and enhancement of the dura of both the brain and spinal cord [6]. Often times this lesion is confused with an inflammatory process such as neurosarcoïdosis [7]. The clinical presentation is also rather nonspecific and can include focal neurological deficits, headaches due to increased intracranial pressure and changes in mental status [1-3,7].

In this report we present the case of 7-year-old boy with diffuse leptomeningeal oligodendrogliomatosis. As with many clinical scenarios of this disease, the initial clinical presentation is nonspecific. The progress of disease is slow with imaging findings developing sometime after the initial clinical presentation. We also report the full pathology and molecular work up of the lesion and make comparisons to the literature describing other clinical presentations of this tumor.

Case Presentation
A 7-year-old boy presented with a five-day history of headaches, vomiting, and gait disturbance. Examination revealed bilateral papilledema and cranial CT showed dilated ventricular system indicative of non-obstructive hydrocephalus. MRI confirmed the CT findings and did not display any brain parenchymal changes or leptomeningeal enhancement. It should be noted that six months prior to admission, a non-contrast enhanced CT scan, performed at another institution for mild head trauma, already showed ventricular dilatation. A Ventriculo-Peritoneal shunt was inserted, and the boy was discharged from hospital five days later. Cytological examination of the CSF was negative. Examination 3 months later, showed that the papilledema had resolved.

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evolution is slow. The initial physical manifestations of this disease are varied, including focal neurological signs of increased intracranial pressure, convulsions and altered mental status. This entity may mimic non-tumoral diseases like neurosarcoidosis or infectious agents that can cause chronic meningitis [7].

The entity was first described by Beck and Russell in 1942 in which they described four cases of “Oligodendrogliomatosis of cerebrospinal pathway” [8]. A series of 36 cases found high proportion of pediatric patients with a male predominance in 66% [2]. Similar to the patient that we presented, cytology form the CSF in this series was negative in 16 patients, indicating that definitive diagnosis could only be obtained by biopsy [2]. However, non-diagnostic brain, pia and arachnoid biopsies are a common phenomena and sometimes several attempts are necessary to arrive at the final diagnosis. As mentioned the CSF samples are typically negative for malignant cells and have a high protein concentration [7]. Lesions of this disease are mainly located in the subarachnoid space; it is unclear whether its origin is in the leptomeningeal compartment or leptomeningeal extension of a subtle intraparenchymal disease [3].

Histologically this lesion shows a classic oligodendrogial pattern: Round cells with surrounding halos and clear cytoplasm. The cells displayed no significant atypia and no mitotic figures were identified. The neoplastic cells were immune positive, for OLIG2 and, S100 protein and also for GFAP and MAP2 (Figure 2). They did not stain for synaptophysin, IDH1-R132H and p53, A diagnosis of disseminated oligodendrogial - like leptomeningeal tumor was made. Molecular analysis demonstrated a BRAF- KIAA1549 fusion, confirming the pathological diagnosis.

Discussion

We present a case of a 7-year-old boy with diffuse leptomeningeal glioneuronal tumor. As with many cases of this disease, the presentation is nonspecific, and the clinical...
shown to have good prognostic outcome while at the same time serving as a possible therapeutic target, however future studies are needed for validation [12,13].

The radiological features of this disease show diffuse leptomeningeal involvement with enhancement throughout the brain and spine on MRI studies [2]. The presence of concomitant hydrocephalus is not uncommon. In the initial stages of disease, the enhancement of the spine is thin, and it looks as the nerves roots are coated with paint [2]. As the disease progresses the enhancement in both the brain and spinal cords shifts from the subarachnoid space to the intra-axial compartment and the lesions be-

Figure 2: Staining of tissue section from biopsy specimen: A) H&E staining; B) Staining for OLIG2; C) Staining for KI67.
come more nodular in nature \[6,14\]. There are reports of cystic lesions in the posterior fossa which are hyperintense on T2 weighted imaging \[2\]. The imaging findings of this lesion are rather nonspecific, and it is difficult to make the diagnosis alone based on radiological findings.

In this study we have presented the case of 7-year-old male patient with diffuse leptomeningeal oligodendrogliomatosis. The initial presenting symptoms were rather nonspecific as were the radiological findings. As we have described this lesion is a complicated entity, which can be mistaken for infectious processes. However, we emphasize to clinicians that leptomeningeal oligodendrogliomatosis of childhood must be kept in the differential diagnosis with patients presenting with nonspecific neurologic findings in combination with diffuse subarachnoid enhancement on brain and spine MRI.

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