A Possible Association between Arnold-Chiari Type I Malformation and Epilepsy: Case Series and Review

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Introduction

Hans Chiari is the one credited with diagnosing Chiari malformations, dating back to 1890 \[1\]. Chiari malformations consists of four types of hindbrain abnormalities with the majority being a type I or type II CM-which is characterized by displacement of part of the cerebellum and brain stem into the foramen magnum, usually accompanied by myelomeningocele \[2\]. In this paper, the authors discuss the incidence and prevalence of epilepsy in a subset of pediatric patients harboring Chiari malformations type I (CM-I).

The displacement of the cerebellar tonsils is a classic hallmark of CM-I. Some authors suggest the definition of a cerebellar tonsil position $\geq 5$ mm below the foramen magnum \[3\]. CM-I is a rare disease characterized by herniation of cerebellar tonsils below foramen magnum with associated anomalies of the posterior fossa \[4\]. It also has a wide array of symptoms, some of which are nonspecific such as headache and neck pain \[3\]. Of notice, 15\%-20\% of CM-I patients may develop hydrocephalus \[5\].

Epileptic seizures have been previously reported in patients with CM-I as an incidental finding \[6-8\]. Yet the widespread use of MRI in the diagnostic work up for children with epilepsy has shown incidental findings of CM-I in some children \[9\], thus de facto increasing the number of patients seen harboring both pathologies.

To date, no agreed-upon management exists in the literature. The authors report on 4 cases of epilepsy in CM-I patients which completely resolved without need for surgical correction of their cerebellar tonsillar herniation as well as a review of existing literature.

Methods

30 pediatric patients at a single institution which underwent MRI of the brain due to either headaches or other reasons and diagnosed with CM-I radiographically were reported. Patients underwent C spine MRIs as well. In patients with suspected epilepsy, electroencephalograms were done demonstrating epileptiform abnormalities. All patients underwent at least one more brain MRI imaging and were followed up by a treating pediatric neurologist.

Results

In our cohort we had a total of 30 patients. 18 were males and 12 were females, the average age was 8.43 years at diagnosis of CM-I. The vast majority of patients in the cohort underwent MRI imaging due to headaches, (10 patients). Some underwent MRI imaging of the brain due to the following: Failure to thrive (1), platybasia (1), precocious puberty (1), photophobia (1), dizziness (1), growth hormone deficiency workup (1) and 4 due to epilepsy. 10 other patients underwent brain imaging due to various unrelated reasons. Tonsillar herniation averaged at 10.2 millimeters upon diagnosis.

The mean follow-up was 22.3 months in all CM-I patients. Serial MRIs showed a consistent CM-I malformation without progression. One patient presented with ventricular dilatation. Only one patient had a cervical spine syrinx on imaging.

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Patients were diagnosed first with an MRI and a follow up including a second MRI was performed at 6 or 12 months as part of a follow up. Timing of the second MRI was determined by the pediatric neurologist taking care of the patient.

Of notice are the following patients without epilepsy in this cohort. One patient had a spontaneous resolution of his Chiari with no tonsillar herniation on a routine follow up imaging, (Figure 1A and Figure 1B), as well as disappearance of the syrinx. This patient presented with headaches and vomiting which resolved during follow up. The imaging which did not show tonsillar herniation was done 4 years after initial diagnosis. One patient had a bilateral choroid plexus papilloma. This patient presented with papilledema which resolved on follow up. He is currently asymptomatic. Another patient had initial diplopia which resolved after 6 weeks. Urinary and fecal incontinence spontaneously resolved in another patient.

**CM-I and Epilepsy**

There were 4 patients who presented with epilepsy. Three patients experienced absence epilepsy and one had focal epilepsy as witnessed by their parents and reported to the treating neurologist on scheduled follow ups. These epileptic fits were later on confirmed with an EEG recording device displaying abnormal activity. All four patients were epilepsy free on follow up (12, 12, 23 and 31 months after initial diagnosis was made), and the median follow-up was 21 months per patient. All 4 patients were put on antiepileptic drugs in which 2 of them epilepsy drugs were stopped during follow up as these patients were seizure free for several months. The seizures did not recur in any of the four patients.

Of the four epilepsy patients, one presented with headaches with epilepsy appearing later during follow up, and 3 presented with an epileptic attack upon presentation. Tonsillar herniation was 9, 6, 6 and 6 millimeters for these 4 patients. None had any other CNS abnormalities (syrinx or ventricular dilatation), aside for one patient with cortical dysplasia seen on brain MRI imaging. These four patients are depicted in Table 1.

**Discussion**

The authors present 4 cases out of a cohort of 30 cases with CM-I which diagnosed with epilepsy between the years 2010-2017 in one institution. Meaning that 13% of the cohort had epilepsy. All patients in this series did not go any neurosurgical treatment but were followed up for a period of 23 months.

There is wide accepted agreement that symptomatic CM-I patients presenting with brainstem dysfunction, cranial nerves deficits, syringomyelia or kyphosis should be operated on. Whereas asymptomatic CM-I patients should not be operated on [10]. This may present as a challenge to the treating physician as he/she must bear in mind that other, coexistent disease might appear with CM-I which will not be amended by surgery alone. Several studies exist describing outcome in CM-I patients undergoing surgery whereas fewer studies exist describing nonsurgical CM-I patients with epilepsy. Buoni, et al. described disappearance of EEG abnormalities after
agree with this view and recommend a watch and wait strategy in CM-I patients presenting with epilepsy after a complete workout containing brain and cervical spine imaging, EEG and close follow up.

**Study Limitations**

In this cohort of 30 patients, patients with CM-I and a report of witnessed epilepsy underwent an EEG study. However not all parents were routinely asked whether or not epilepsy symptoms have occurred in their children. In addition, not all patients underwent an EEG as part of their work up for CM-I, which may have resulted in undetected instances of epilepsy in some patients and an underestimation of the actual number of children with CM-I malformations and epilepsy. In addition, the four children diagnosed with CM-I and epilepsy were diagnosed at a smaller age than the rest of the group. This may lead to the assumption that patients with epilepsy and CM-I are diagnosed at an earlier age. However, this number is too small to assume this assumption.

**Conclusion**

Epilepsy in CM-I is an unexplained entity. In this cohort of patients however, it appeared in higher frequency in patients harboring CM-I pathologies. Our study raises the possibility of the dual occurrence of epilepsy and CM-I in which epilepsy is transient after several years. The use of anti-epileptic drugs which are eventually tapered down to a minimum or stopped altogether is an option in this group of patients. The medical staff treating these patients must bear in mind that CM-I and epilepsy can coexist albeit for a short period of time in a CM-I patient’s life when taking care of these patients. Further studies are needed in order to clarify the associ-
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