Chiari 1 Malformation and Epilepsy in Children: A Missing Relationship

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Abstract: Purpose: Once believed a result of pathophysiological correlations, the association between Chiari 1 malformation (CM1) and epilepsy has since been considered as a coincidence, due to missing etiologic or clinical matching points. At present, the problem is being newly debated because of the increasing number of CM1 diagnoses, often among children with seizures. No specific studies on this topic are available yet. The present study aimed at updating the information on this topic by reporting on a series of children specifically enrolled and retrospectively analyzed for this purpose. Methods: All children admitted between January 2015 and June 2020 for epilepsy and CM1 were considered (Group 1). They were compared with children admitted in the same period for symptoms/signs related to CM1 and/or syringomyelia (Group 2). Syndromic patients were excluded, as well as those with tumoral or other overt intracranial lesions. All patients received a complete preoperative work-up, including MRI and/or syringomyelia (Group 2). Symptomatic children with CM1/syringomyelia were operated on. The pertinent literature was reviewed. Results: Group 1 was composed of 29 children (mean age: 6.2 years) showing CM1 and epilepsy with several types of seizures. A share of 27% had CM1-related symptoms and syringomyelia. The mean tonsillar ectopia was 7.5 mm. Surgery was performed in 31% of cases. Overall, 62% of children are currently seizure-free (including 5/9 children who were operated on). Tonsillar herniation and syringomyelia regressed in 4/9 cases and 4/8 cases, improved in 4/9 cases and 3/8 cases, and remained stable in 1/9 and 1/8 cases, respectively. CM1 signs/symptoms regressed completely in 6/8 cases and improved or remained stable in one case in each of the two remaining patients. Group 2 consisted of 77 children (mean age: 8.9 years) showing symptoms of CM1 (75%) and/or syringomyelia (39%). The mean tonsillar ectopia was 11.8 mm. Non-specific EEG anomalies were detected in 13 children (17%). Surgery was performed in 76.5% of cases (18 children were not operated on because of oligosymptomatic). Preoperative symptoms regressed in 26%, improved in 50%, remained stable 22%, and worsened in 2%; CM1 radiologically regressed in 39%, improved in 37%, remained unchanged in 22%, and worsened in 2%; and syringomyelia/hydromyelia regressed in 61%, improved in 30%, and was stable in 9%. No statistically significant differences between the two groups were detected regarding the M/F ratio, presence of syringomyelia/hydromyelia, or CM1/syringomyelia outcome; moreover, no correlation occurred between seizure-free condition and PF decompression in Group 1, or between disappearance of EEG anomalies and PF decompression in Group 2. A significant difference between the two groups was noticed regarding the mean age at admission ($p = 0.003$), amount of tonsillar herniation ($p < 0.00001$), and PF decompression ($p = 0.0001$). Conclusions: These findings do not support clinical correlations between CM1 and epilepsy. Their course depends on surgery and antiepileptic drugs, respectively. The analysis of the literature does not provide evidence of a relationship between seizures and cerebellar anomalies such as CM1. Rather than being linked to a syndrome that could explain such an association, the connection between the two now has to be considered to be random.

Keywords: Chiari 1 malformation; syringomyelia; epilepsy; seizures; posterior fossa; EEG; cerebellum
1. Introduction

The association between Chiari type I malformation (CM1) and epilepsy raises relevant interest among clinicians and researchers. The first reason for such an interest is related to the growing number of new diagnoses of CM1, both in children and adult [1,2]. The increasing numbers of patients with diagnosed CM1 (and syringomyelia) require increasing efforts for their correct classification and treatment, a need that is exacerbated by the still-missing guidelines about CM1 management [3]. One of the results of the greater “attention” paid to CM1 patients is their division into different subtypes [4], of which the CM1/epilepsy groups are potentially one. On the other hand, epilepsy is already known to be a common problem, with a 0.5–1% prevalence in the general population [5] and 8% mean prevalence in hospitalized patients worldwide [6]. Therefore, its impact on the population’s health is significant.

On these grounds, the association between the previous two conditions is regarded with concern because of the known impact of epilepsy in the clinical practice and the potential role of CM1 in further increasing that impact. A peculiar issue, which has been addressed in the past, regards the possible etiological and clinical relationships between the two conditions. No conclusions have been reached so far about whether this association may be considered to be random. The present study aims to update the information on this topic by reporting on a series of children specifically enrolled and analyzed for this purpose and by reviewing the pertinent literature.

2. Materials and Methods

All children consecutively admitted between January 2015 and June 2020 for epilepsy and harboring CM1 malformation were considered for the present study. They were compared with children admitted in the same period for symptoms/signs related to CM1 and/or syringomyelia. The aforementioned time period was considered to include patients with homogeneous preoperative and postoperative evaluation, and to have a minimum follow-up of 2 years.

Epilepsy and seizures were classified according to ILAE 2017 classification [7]. Epilepsy was managed by antiepileptic drugs according to the patients’ characteristics. CM1 was radiologically defined as tonsillar caudal ectopia ≥ 5 mm, associated with peg-like appearance of the tonsils or other signs of posterior fossa (PF) overcrowding (e.g., effacement of CSF spaces) [8]. Surgery was performed in clearly symptomatic patients and/or with large or progressing syringomyelia [3]. PF decompression was performed to manage CM1, by suboccipital craniectomy and C1 laminectomy, with/without dural opening (and tonsil shrinkage) based on the patient’s characteristics. Endoscopic third ventriculostomy was used to manage associated hydrocephalus.

All children received preoperative and postoperative neurological examination, neuropsychological tests, brain (with perfusion sequences) and whole spinal cord MRI, and surface EEG. In selected cases, accessory investigations, such as motor and/or somatosensory potentials and polysomnography, were performed (their results are not discussed here since their impact was not relevant). Syndromic patients were excluded, as well as those with tumoral or other overt intracranial lesions (e.g., arachnoid cysts), incomplete preoperative/postoperative work-up, or details lost at follow-up.

Patients with epilepsy and CM1 were mainly referred to the Pediatric Neurology department of our institution, while those with CM1 were referred to the Pediatric Neurosurgery department. The first subset of patients was included in Group 1 of the present study, while those with CM1 belonged to Group 2. The mean follow-up was 40.68 months (range: 24–66 months).

The following variables were considered for the comparison between the two groups: age, sex, presence of EEG anomalies, cerebellar tonsils herniation, presence of hydromyelia/syringomyelia, PF surgical decompression, postoperative regression/improvement of CM1 symptoms, and radiological picture. The statistical analysis was realized through $\chi^2$ test and 2-tailed $t$-test. A $p$ value < 0.05 was considered as statistically significant.
3. Results

3.1. Patients' Characteristics

Overall, 106 children were enrolled for the present analysis. All of them were alive at current follow-up.

Group 1 was composed of 29 children (14 boys, 15 girls) with a 6.2-year mean age at admission (range: 11 months–15 years). All these patients were admitted and studied because of epileptic seizures and all of them were found to have CM1. These children represent 12% of all pediatric patients admitted to our institution for epilepsy in the considered period and 18% of all CM1 cases (both symptomatic and incidentally diagnosed) observed in the same period (157 cases). Regarding seizures, 4 children (13%) presented absences (A), 8 (27.5%) focal motor seizures (FM), 6 (20.5%) focal non-motor seizures (FNM), 4 (13%) focal to generalized tonic-clonic seizures (FGTC), 2 (10%) generalized tonic-clonic seizures (GTC) (Figure 1), 4 (13%) generalized tonic seizures (GT), and 1 (3%) myoclonic seizure (M). All children underwent antiepileptic drug treatment. Eight out of 29 children had signs/symptoms related to CM1 (27%). The mean tonsillar ectopia was 7.5 mm (range: 5–13 mm) (Figure 2). Eight patients (27%) harbored syringomyelia (5 cases) or hydromyelia (3 cases), and 2 cases were associated with hydrocephalus. No other relevant anomalies or microlesions were detected on MRI. The details of Group 1 patients are reported in Table 1.

Figure 1. Six-year-old boy with C1M and GTC: (A) interictal diffuse spike waves during sleep; (B) interictal focal spike waves (fronto-central regions) during sleep.
Figure 1. Six-year-old boy with C1M and GTC: (A) interictal diffuse spike waves during sleep; (B) interictal focal spike waves (fronto-central regions) during sleep.

Figure 2. 3T MRI of the same case in Figure 1. On sagittal view, C1M is evident, with moderate hypoplasia of PF and 8 mm ectopia of cerebellar tonsils (A, arrow). No relevant findings are visible on sagittal view after gadolinium administration (B) or on T2 axial views (C, D).

Table 1. Clinical characteristics of Group 1.

| Patients (No and Sex) | Age (Years) | Seizures | Interictal EEG | Developmental Problems | Tonsils Ectopia (mm) | Spinal Cord and Other Findings | C1M/Syringomyelia Symptoms |
|-----------------------|-------------|----------|----------------|------------------------|----------------------|-------------------------------|---------------------------|
| 1, F                  | 7           | FM       | bilateral S and SW | No                     | 9                    | Normal                        | no                        |
| 2, F                  | 1           | GT       | bouffées of slow W with S (parieto-occipital region bilaterally) | No                     | 13                   | cervico-thoracic syringomyelia | cough, headache, ataxia    |
| 3, F                  | 5           | GTC      | frequent S, PS and SW (temporal and parieto-occipital region bilaterally) | Autism              | 6                    | hydromyelia (ventriculomegaly) | cough, headache, ataxia, dysphagia |
| 4, M                  | 6           | FGTC     | isolated or couple of S and SW (right central and temporal regions) | behavioral disturbance | 12                   | Normal                        | no                        |
| 5, F                  | 5           | A        | frequent, bilateral SW and PSW | No                     | 6                    | Normal                        | no                        |
| 6, M                  | 15          | GT       | left hemispheric S and SW | Autism              | 10                   | cervico-thoracic syringomyelia | no                        |
| 7, M                  | 12          | GTC      | fronto-central epileptiform activity with generalization | language delay | 8                    | normal                        | no                        |
| 8, F                  | 6           | FM       | right posterior regions PSW | psychomotor delay | 6                    | cervico-thoracic syringomyelia | central apneas             |
| 9, M                  | 4           | A        | PSW | No                     | 5                    | normal                        | no                        |
| 10, M                 | 2           | A        | right occipital epileptiform activity | psychomotor delay | 6                    | hydromyelia (ventriculomegaly) | cough, headache            |
| 11, M                 | 1           | FM       | focal right fronto-central epileptiform activity | language delay | 10                   | normal                        | no                        |
| 12, M                 | 6           | FNM      | bilateral PSW | No                     | 5                    | normal                        | no                        |
| 13, F                 | 7           | M        | slow W with S and myoclonic activity | behavioral disturbance | 13                   | normal                        | cough, headache            |
| 14, F                 | 9           | GT       | left temporal and posterior regions slow W and slow SW | No                     | 6                    | syringomyelia (ventriculomegaly) | cough, headache, dizziness |
### Table 1. Cont.

| Patients (No and Sex) | Age (Years) | Seizures | Interictal EEG | Developmental Problems | Tonsils Ectopia (mm) | Spinal Cord and Other Findings | CIM/Syringomyelia Symptoms |
|----------------------|-------------|----------|----------------|------------------------|----------------------|-------------------------------|-----------------------------|
| 15, M 7 FNM/SD       | 7           | FNM      | right frontal SW | psychomotor delay       | 5                    | normal                        | cough, headache, lower limbs paresthesia |
| 16, F 8 A            | 8           | A        | fronto-central sharp W | behavioral disturbance | 5                    | normal                        | no                          |
| 17, F 7 FNM/SD       | 7           | FNM      | right posterior regions slow W | No | 5                    | normal                        | neck pain, upper and lower limbs paresthesia |
| 18, F 6 FNM/SD       | 6           | FM       | left frontal SW | No                     | 8.5                  | cervico-thoracic syringomyelia | cough, headache, neck pain |
| 19, M 4 GT           | 4           | GT       | bilateral epileptiform activity | No | 6                    | normal                        | cough, headache             |
| 20, F 5 FM           | 5           | FM       | right fronto-central sharp W | No | 5                    | normal                        | no                          |
| 21, F 6 FNM/SD       | 6           | FNM      | left temporal slow W | No | 6.5                  | normal                        | no                          |
| 22, M 4 FM           | 4           | FM       | right central SW and theta activity | No | 10                  | normal                        | cough, headache, upper limbs paresthesia |
| 23, M 8 FGTC/SD      | 8           | FGTC     | posterior regions biphasic S and sharp W | No | 11                  | cervical syringomyelia       | cough, headache, neck pain |
| 24, F 7 FGTC/SD      | 7           | FGTC     | right frontal and temporal SW | psychomotor delay       | 6                    | normal                        | no                          |
| 25, M 8 FM           | 8           | FM       | left centro-temporal slow W and theta activity | No | 5                    | normal                        | no                          |
| 26, M 5 FGTC/SD      | 5           | FGTC     | right posterior regions slow W and S | Autism | 5                   | normal                        | no                          |
| 27, M 8 FNM/SD       | 8           | FNM      | right centro-temporal slow W | behavioral disturbance | 9                    | normal                        | no                          |
| 28, F 3 FM           | 3           | FM       | right central slow W | No | 10                  | normal                        | no                          |
| 29, F 10 FNM/SD      | 10          | FNM      | left fronto-temporal S and PS | No | 6                    | normal                        | no                          |

A: absences, FM: focal motor seizures, FNM: focal non-motor seizures, FGTC: focal to generalized tonic-clonic seizures, GTC: generalized tonic-clonic seizures, GT: generalized tonic seizures, M: myoclonic seizures. PS: poly-spike, PSW: poly-spike-waves, S: spikes, SW: spike-waves, W: waves.

Group 2 consisted of 77 children (43 boys, 34 girls) with an 8.9-year mean age at admission (range: 3–18 years). All these children were admitted because of CM1 with/without syringomyelia. All of them were symptomatic, but signs/symptoms clearly related to CM1/syringomyelia were present in 75% of cases (58 children) (Figure 3). Occipital cough and headache were the main symptoms, followed by dizziness, nystagmus, ataxia, paresthesia, and dysphagia. The mean tonsillar ectopia was 11.8 mm (range: 5–28 mm). Thirty patients (39%) harbored syringomyelia (21 cases) or hydromyelia (9 cases), and 3 cases were associated with ventriculomegaly. No other relevant anomalies or microlesions were detected on MRI. EEG anomalies were detected in 13 children (17%), being represented by slow waves in 6 cases (Figure 4), theta activity in 3 cases, sporadic spikes in 2 cases, and sharp waves, spikes-waves, and poli-spikes in the remaining 2 cases. None of these
13 children showed epileptic seizures nor received antiepileptic drugs. The details on Group 2 patients are reported on Table 2.

Figure 3. Four-year-old girl with symptomatic C1M. (A) MRI, sagittal view, showing a PF moderate hypoplasia with peg-like appearance of the cerebellar tonsils that present a 15 mm downward herniation below the McRae line (arrow). Note the disappearance of the CSF flow through the posterior subarachnoid spaces on cine-MRI (B, double arrows). No relevant findings are found on brain axial view (C), while a cervico-thoracic syringomyelia is evident on spinal cord MRI (D, arrow).

Table 2. Characteristics of Group 2.

| Patients No | Sex | Age (Years) | Signs/Symptoms | Tonsils Ectopia (mm) | Spinal Cord and Other Findings | EEG Anomalies |
|-------------|-----|-------------|----------------|---------------------|-------------------------------|---------------|
| 1           | M   | 7           | psychomotor delay, dizziness, ataxia | 8                   | cervico-thoracic hydromyelia | theta activity |
| 2           | F   | 14          | headache, dizziness, ataxia, dysphagia | 11                  | normal                         |               |
| 3           | F   | 10          | headache, dysphagia, dizziness          | 12                  | cervical hydromyelia           |               |
| 4           | M   | 5           | headache, upper limbs paresthesia       | 20                  | normal                         |               |
| 5           | F   | 16          | cough, headache, upper limbs paresthesia | 15                  | cervico-thoracic syringomyelia |               |
| 6           | M   | 10          | None                              | 5                   | normal                         |               |
| 7           | F   | 15          | headache                          | 15                  | normal                         |               |
| 8           | M   | 6           | None                              | 14                  | cervical hydromyelia           |               |
| Patients No | Sex | Age (Years) | Signs/Symptoms | Tonsils Ectopia (mm) | Spinal Cord and Other Findings | EEG Anomalies |
|------------|-----|-------------|----------------|---------------------|--------------------------------|---------------|
| 9          | F   | 5           | psychomotor delay | 13                  | normal                        | slow waves    |
| 10         | M   | 5           | psychomotor delay | 11                  | cervico-thoracic hydromyelia | slow waves    |
| 11         | F   | 7           | psychomotor delay | 13                  | ventriculomegaly              | /             |
| 12         | M   | 7           | headache, ataxia, dysphagia | 10      | normal                        | slow waves    |
| 13         | F   | 6           | headache, neck pain | 7                   | normal                        | /             |
| 14         | M   | 3           | Ataxia           | 10                  | normal                        | /             |
| 15         | M   | 6           | left hemiparesis | 15                  | cervical syringomyelia        | /             |
| 16         | M   | 11          | cough, headache upper and lower limbs paresthesia | 10      | cervico-thoracic syringomyelia | /             |
| 17         | F   | 3           | cough, headache  | 7                   | normal                        | /             |
| 18         | F   | 4           | cough, headache, psychomotor delay | 11      | cervical hydromyelia         | sharp waves, spike-waves, and poly-spikes |
| 19         | M   | 5           | cough, headache, dizziness | 11      | normal                        | /             |
| 20         | F   | 12          | cough, headache, drop attacks | 20      | normal                        | /             |
| 21         | M   | 7           | cough, headache, psychomotor delay | 9       | normal                        | /             |
| 22         | M   | 13          | ataxia, dysphagia, dizziness, snoring | 10      | cervico-thoracic syringomyelia | /             |
| 23         | F   | 10          | dypleopia, nystagmus, | 12      | cervico-thoracic syringomyelia | /             |
| 24         | M   | 4           | None             | 5                   | normal                        | /             |
| 25         | F   | 6           | dizziness        | 5                   | normal                        | isolated spikes |
| 26         | M   | 18          | hand tremors     | 19                  | cervico-thoracic syringomyelia | /             |
| 27         | F   | 7           | cough, headache  | 18                  | normal                        | /             |
| 28         | M   | 17          | headache, dizziness | 8       | normal                        | /             |
| 29         | F   | 6           | cough, headache  | 21                  | normal                        | /             |
| 30         | M   | 14          | nystagmus, dysphagia | 10      | syringobulbia                 | /             |
| 31         | M   | 6           | dypleopia, right VI cranial nerve palsy | 16      | thoracic syringomyelia       | /             |
| 32         | F   | 4           | ataxia, dysphagia, | 12      | normal                        | /             |
| 33         | F   | 18          | neck pain, central sleep apneas | 8       | normal                        | /             |
| 34         | M   | 6           | cough, headache  | 7                   | normal                        | /             |
| 35         | F   | 8           | Nystagmus        | 9                   | cervical hydromyelia         | /             |
| 36         | M   | 16          | nystagmus, ataxia | 20      | normal                        | /             |
| 37         | F   | 18          | cough, headache, dizziness | 15      | normal                        | /             |
| 38         | M   | 10          | headache, dizziness | 8       | cervical hydromyelia         | /             |
| 39         | F   | 15          | cough, headache, nystagmus | 9       | cervico-thoracic syringomyelia | theta activity |
Table 2. Cont.

| Patients No | Sex | Age (Years) | Signs/Symptoms | Tonsils Ectopia (mm) | Spinal Cord and Other Findings | EEG Anomalies |
|-------------|-----|-------------|----------------|----------------------|-------------------------------|---------------|
| 40          | F   | 4           | cough, headache, dizziness | 15                   | cervico-thoracic hydromyelia | slow waves    |
| 41          | M   | 13          | Headache        | 7                    | normal                         |               |
| 42          | F   | 7           | right lower limb pain | 5                    | normal                         |               |
| 43          | F   | 7           | psychomotor delay | 9                    | cervico-thoracic syringomyelia | isolated spikes |
| 44          | M   | 5           | None            | 7                    | normal                         |               |
| 45          | M   | 17          | cough, headache | 14                   | normal                         |               |
| 46          | M   | 18          | headache, dizziness, lower limbs weakness | 9 | normal |               |
| 47          | F   | 10          | dizziness       | 5                    | normal                         | sharp waves and spikes |
| 48          | F   | 6           | dizziness       | 8                    | cervico-thoracic syringomyelia |               |
| 49          | M   | 6           | headache, drop attacks, vomiting | 9 | normal | slow waves |
| 50          | M   | 7           | cough, headache, salorrhoea | 12 | normal |               |
| 51          | F   | 12          | Scoliosis       | 28                   | holocord syringomyelia        |               |
| 52          | M   | 13          | Headache        | 7                    | normal                         |               |
| 53          | F   | 5           | headache, dizziness | 11                   | normal                         |               |
| 54          | M   | 5           | headache, dizziness | 11                   | cervico-thoracic syringomyelia |               |
| 55          | M   | 8           | right hypoacusis | 9                    | normal                         |               |
| 56          | F   | 2           | cough, headache, upper limbs paresthesia | 17 | normal |               |
| 57          | M   | 3           | cough, headache, central sleep apneas | 5 | normal |               |
| 58          | F   | 8           | cough, headache, vomiting | 8 | cervico-thoracic hydromyelia |               |
| 59          | M   | 3           | psychomotor delay | 9                    | normal                         |               |
| 60          | M   | 15          | headache, neck pain | 12                   | ventriculomegaly               |               |
| 61          | M   | 10          | headache        | 10                   | normal                         |               |
| 62          | M   | 8           | cough, headache | 10                   | normal                         |               |
| 63          | F   | 7           | headache, upper and lower limbs paresthesia | 31 | cervical syringomyelia |               |
| 64          | F   | 6           | headache, enuresis | 9                    | thoracic syringomyelia        |               |
| 65          | M   | 14          | cough, headache, lower limbs paresthesia | 9 | thoracic syringomyelia |               |
| 66          | F   | 6           | Nystagmus       | 6                    | normal                         |               |
| 67          | M   | 9           | cough, headache, lower limbs pain | 21 | cervico-thoracic syringomyelia |               |
| 68          | M   | 6           | headache, febrile paroxysms | 12 | normal | theta activity |
Table 2. Cont.

| Patients No | Sex | Age (Years) | Signs/Symptoms       | Tonsils Ectopia (mm) | Spinal Cord and Other Findings                       | EEG Anomalies |
|-------------|-----|-------------|----------------------|----------------------|------------------------------------------------------|---------------|
| 69          | F   | 15          | headache, dizziness  | 18                   | cervico-thoracic syringomyelia                       | /             |
| 70          | F   | 7           | Scoliosis            | 12                   | normal                                               | /             |
| 71          | M   | 16          | central sleep apneas | 18                   | normal                                               | slow waves    |
| 72          | F   | 5           | cough, headache      | 13                   | normal                                               | /             |
| 73          | F   | 12          | cough, headache      | 15                   | cervico-thoracic hydromyelia                        | /             |
| 74          | M   | 9           | headache, vomiting   | 8                    | ventriculomegaly                                    | /             |
| 75          | M   | 9           | headache, neck pain, ataxia | 7               | cervical syringomyelia                              | /             |
| 76          | M   | 7           | ataxia, drop attacks | 11                   | normal                                               | /             |
| 77          | M   | 9           | cough, headache, dizziness | 22              | cervical syringomyelia                              | /             |

3.2. Treatment and Outcome

In Group 1, surgery (for CM1 and/or ventriculomegaly) was performed in 9 cases (31%): PF decompression was realized in 8/9 cases (in one patient following ETV), while ETV alone was performed in the remaining patient. Five of the 9 children operated on are seizure-free. Overall, 62% of children are currently seizure-free (18 cases). On neuroimaging, tonsillar herniation and syringomyelia/hydromyelia regressed at follow-up in 4/9 cases and 4/8 cases, improved in 4/9 cases (including the patient who received only ETV) and 3/8 cases, and remained stable in 1/9 and 1/8 cases, respectively. CM1 signs/symptoms regressed completely in 6/8 cases (one patient was asymptomatic but operated on for large syringomyelia, one underwent ETV alone) and improved or remained stable in one case each of the two remaining patients. Among the 21 non-operated-on cases, the PF radiological picture remained substantially unchanged, except for 2 cases where the tonsillar herniation spontaneously improved (Table 3).

In Group 2, surgery was performed in 59 cases (76.5%): PF decompression was realized in 54/59 cases because of symptomatic CM1 and/or large or progressing syrinx, while the remaining 5 patients received ETV for hydrocephalus (3 cases) and invasive recording of intracranial pressure (ICP), which ruled out raised ICP. The remaining 18 children were not operated on because they were oligosymptomatic. At follow-up, 15 out of these 18 children had their radiological picture unchanged, while a spontaneous improvement in CM1 was detected in the remaining 3. All 3 patients undergoing ETV had normalization of hydrocephalus and improvement in their CM1/syringomyelia-related clinical and radiological picture. Finally, among patients operated on for CM1/syringomyelia: preoperative symptoms regressed in 14 cases (26%), improved in 27 (50%), remained stable in 12 (22%), and worsened in one (2%); on neuroimaging, CM1 radiologically regressed in 21/54 children (39%), improved in 20/54 (37%), remained unchanged in 12/54 (22%), and worsened in 1/54 (2%); and in syringomyelia/hydromyelia, regressed in 14/23 children (61%), improved in 7/23 (30%), and was stable in 2/23 (9%) (Table 4).
### Table 3. Outcome of Group 1.

| Patients (No) | Age (Years) | Seizures | Tonsils Ectopia (mm) | Spinal Cord and Other Findings | C1M/Syringomyelia Symptoms | Surgery | Epilepsy Outcome | C1M/Syrinx Outcome |
|---------------|-------------|----------|----------------------|-----------------------------|--------------------------|---------|-----------------|-------------------|
| 1             | 7           | FM       | 9                    | normal                       | No                       | none    | seizure-free    | Stable            |
| 2             | 1           | GT       | 13                   | cervico-dorsal syringomyelia | cough headache, ataxia   | PF decompression + tonsils coagulation | seizure-free | C1M, syrinx and headache regressed, ataxia improved |
| 3             | 5           | GTC      | 6                    | hydromyelia (ventriculomegaly) | cough headache, ataxia, dysphagia | ETV; bony PF decompression + C1 laminectomy | refractory seizures | C1M and symptoms regressed, hydromyelia improved |
| 4             | 6           | FGTC     | 12                   | normal                       | No                       | None    | >90% reduction  | Stable            |
| 5             | 5           | A        | 6                    | normal                       | no                       | none    | seizure-free    | Stable            |
| 6             | 15          | GT       | 10                   | cervico-dorsal syringomyelia | no                       | bony PF decompression + C1 laminectomy | seizure-free | C1M and syrinx stable |
| 7             | 12          | GTC      | 8                    | normal                       | no                       | none    | seizure-free    | Stable            |
| 8             | 6           | FM       | 6                    | cervico-dorsal syringomyelia | central apneas           | PF decompression + duraplasty | >90% reduction | C1M improved, syrinx and symptoms regressed |
| 9             | 4           | A        | 5                    | normal                       | No                       | none    | >90% reduction  | Stable            |
| 10            | 2           | A        | 6                    | hydromyelia                  | cough headache           | bony PF decompression + C1 laminectomy | >90% reduction | C1M, hydromyelia and symptoms regressed |
| 11            | 1           | FM       | 10                   | normal                       | No                       | none    | seizure-free    | Stable            |
| 12            | 6           | FNM      | 5                    | normal                       | No                       | none    | seizure-free    | Stable            |
| 13            | 7           | M        | 13                   | normal                       | cough headache           | none    | >90% reduction  | Stable            |
| 14            | 9           | GT       | 6                    | syringomyelia (ventriculomegaly) | cough headache, vertigo | ETV    | seizure-free    | C1M and syrinx improved, symptoms regressed |
| 15            | 7           | FNM      | 5                    | normal                       | cough headache, lower limbs paresthesia | none | >90% reduction | Stable            |
| 16            | 8           | A        | 5                    | normal                       | No                       | none    | seizure-free    | Stable            |
| 17            | 7           | FNM      | 5                    | normal                       | neck pain, upper and lower limbs paresthesia | none | refractory seizures | Stable            |
| 18            | 6           | FM       | 8.5                  | cervico-dorsal syringomyelia | cough headache, neck pain | PF decompression + duraplasty | seizure-free | C1M improved, syrinx regressed, symptoms stable |
| 19            | 4           | GT       | 6                    | normal                       | cough headache           | bony PF decompression + C1 laminectomy | seizure-free | C1M improved, symptoms regressed |
| 20            | 5           | FM       | 5                    | normal                       | No                       | none    | >90% reduction  | Stable            |
| 21            | 6           | FNM      | 6.5                  | normal                       | no                       | none    | seizure-free    | Stable            |
| 22            | 4           | FM       | 10                   | normal                       | cough headache, upper limbs paresthesia | none | seizure-free | C1M and symptoms spontaneously improved |
| 23            | 8           | FGTC     | 11                   | cervical syringomyelia       | cough headache, neck pain | bony PF decompression + C1 laminectomy | >90% reduction | C1M and symptoms regressed, syrinx improved |
| 24            | 7           | FGTC     | 6                    | normal                       | No                       | none    | seizure-free    | Stable            |
| 25            | 8           | FM       | 5                    | normal                       | No                       | none    | seizure-free    | Stable            |
| 26            | 5           | FGTC     | 5                    | normal                       | no                       | none    | >90% reduction  | Stable            |
| 27            | 8           | FNM      | 9                    | normal                       | No                       | none    | seizure-free    | C1M spontaneously improved |
| 28            | 3           | FM       | 10                   | normal                       | no                       | none    | seizure-free    | Stable            |
| 29            | 10          | FNM      | 6                    | normal                       | no                       | none    | seizure-free    | Stable            |

A: absences; ETV: endoscopic third ventriculostomy; FM: focal motor seizures; FNM: focal non-motor seizures; FGTC: focal to generalized tonic-clonic seizures; GTC: generalized tonic-clonic seizures; GT: generalized tonic seizures; M: myoclonic seizures; PF: posterior fossa.
Table 4. Outcome of Group 2.

| Patients No | Age (Years) | Tonsils Ectopia (mm) | Spinal Cord and Other Findings | EEG Anomalies | Surgery | CIM/Syrinx Outcome | EEG Anomalies Outcome |
|-------------|-------------|----------------------|--------------------------------|---------------|---------|-------------------|----------------------|
| 1           | 7           | 8                    | cervico-thoracic hydromyelia  | theta activity | bony PF decompression + C1 laminectomy | CIM and syrinx improved, symptoms stable | unchanged            |
| 2           | 14          | 11                   | normal                         | /             | bony PF decompression + C1 laminectomy | CIM regressed, symptoms improved | /                    |
| 3           | 10          | 12                   | cervical hydromyelia           | /             | bony PF decompression + C1 laminectomy | CIM stable, syrinx regressed, symptoms improved | /                    |
| 4           | 5           | 20                   | normal                         | /             | bony PF decompression + C1 laminectomy | CIM stable, symptoms improved | /                    |
| 5           | 16          | 15                   | cervico-thoracic syringomyelia | PF decompression + tonsils coagulation | PF decompression + tonsils coagulation | CIM and syringomyelia regressed, symptoms stable | /                    |
| 6           | 10          | 5                    | normal                         | /             | none                              | CIM unchanged           | /                    |
| 7           | 15          | 15                   | normal                         | /             | none                              | CIM unchanged           | /                    |
| 8           | 6           | 14                   | cervical hydromyelia           | /             | ICP monitoring                     | CIM and syringomyelia unchanged | /                    |
| 9           | 5           | 13                   | normal                         | slow waves    | none                              | CIM unchanged           | unchanged            |
| 10          | 5           | 11                   | cervico-thoracic hydromyelia  | slow waves    | none                              | CIM and syringomyelia unchanged | unchanged            |
| 11          | 7           | 13                   | ventriculomegaly               | /             | bony PF decompression + C1 laminectomy | CIM unchanged, symptoms improved | /                    |
| 12          | 7           | 10                   | normal                         | slow waves    | bony PF decompression + C1 laminectomy | CIM and symptoms regressed | unchanged            |
| 13          | 6           | 7                    | normal                         | /             | none                              | CIM unchanged           | /                    |
| 14          | 3           | 10                   | normal                         | /             | none                              | CIM unchanged           | /                    |
| 15          | 6           | 15                   | cervical syringomyelia         | /             | bony PF decompression + C1 laminectomy | CIM regressed, hydromyelia improved, symptoms improved | /                    |
| 16          | 11          | 10                   | cervico-thoracic syringomyelia | PF decompression + tonsils coagulation | PF decompression + tonsils coagulation | CIM, syringomyelia and symptoms regressed | /                    |
| 17          | 3           | 7                    | normal                         | /             | none                              | CIM unchanged           | /                    |
| 18          | 4           | 11                   | cervical hydromyelia           | sharp waves, spike-waves, and poly-spikes | bony PF decompression + C1 laminectomy | CIM, syringomyelia and symptoms regressed | unchanged            |
| 19          | 5           | 11                   | normal                         | /             | bony PF decompression + C1 laminectomy | CIM and symptoms regressed | /                    |
| 20          | 12          | 20                   | normal                         | /             | PF decompression + duraplasty      | CIM regressed, symptoms improved | /                    |
| 21          | 7           | 9                    | normal                         | /             | bony PF decompression + C1 laminectomy | CIM and symptoms improved | /                    |
| 22          | 13          | 10                   | cervico-thoracic syringomyelia | /             | bony PF decompression + C1 laminectomy + C0-C3 fixation | CIM improved, syringomyelia and symptoms regressed | /                    |
| 23          | 10          | 12                   | cervico-thoracic syringomyelia | /             | PF decompression + tonsils coagulation | CIM regressed, symptoms improved | /                    |
| 24          | 4           | 5                    | normal                         | /             | none                              | CIM spontaneously improved | /                    |
| 25          | 6           | 5                    | normal                         | isolated spikes | none                              | CIM unchanged           | disappeared          |
| 26          | 18          | 19                   | cervico-thoracic syringomyelia | /             | PF decompression + tonsils coagulation | CIM and syringomyelia regressed, symptoms improved | /                    |
| 27          | 7           | 18                   | normal                         | /             | PF decompression + tonsils coagulation | CIM regressed, symptoms stable | /                    |
| 28          | 17          | 8                    | normal                         | /             | PF decompression + tonsils coagulation | CIM and symptoms stable | /                    |
| 29          | 6           | 21                   | normal                         | /             | bony PF decompression + C1 laminectomy | CIM and symptoms improved | /                    |
| 30          | 14          | 10                   | syringobulbia                  | /             | bony PF decompression + C1 laminectomy | CIM and syringobulbia regressed, symptoms improved | /                    |
| 31          | 6           | 16                   | thoracic syringomyelia         | /             | PF decompression + tonsils coagulation | CIM, syringomyelia and symptoms regressed | /                    |
### Table 4. Cont.

| Patients No | Age (Years) | Tonsils Ectopia (mm) | Spinal Cord and Other Findings | EEG Anomalies | Surgery | CIM/Syrinx Outcome | EEG Anomalies Outcome |
|-------------|-------------|----------------------|--------------------------------|---------------|---------|--------------------|------------------------|
| 32          | 4           | 12                   | normal                         | /             | bony PF decompression + C1 laminectomy | CIM regressed, symptoms improved | /                      |
| 33          | 18          | 8                    | normal                         | /             | bony PF decompression + C1 laminectomy | CIM and symptoms regressed | /                      |
| 34          | 6           | 7                    | normal                         | /             | none                             | CIM unchanged            | /                      |
| 35          | 8           | 9                    | cervical hydromyelia          | /             | none                             | CIM unchanged            | /                      |
| 36          | 16          | 20                   | normal                         | /             | bony PF decompression + C1 laminectomy | CIM regressed, symptoms improved | /                      |
| 37          | 18          | 15                   | normal                         | /             | bony PF decompression + C1 laminectomy | CIM and symptoms stable | /                      |
| 38          | 10          | 8                    | cervical hydromyelia          | /             | bony PF decompression + C1 laminectomy | CIM and syrinx, symptoms improved | /                      |
| 39          | 4           | 9                    | cervico-thoracic syringomyelia | theta activity | bony PF decompression + C1 laminectomy | CIM, syrinx and symptoms improved | unchanged |
| 40          | 15          | 4                    | cervico-thoracic hydromyelia  | slow waves    | bony PF decompression + C1 laminectomy | CIM and syrinx, symptoms improved | disappeared |
| 41          | 13          | 7                    | normal                         | /             | none                             | CIM spontaneously regressed | /                      |
| 42          | 7           | 5                    | normal                         | /             | none                             | CIM unchanged            | /                      |
| 43          | 7           | 9                    | cervico-thoracic syringomyelia | isolated spikes | 16 Sub + PF decompression + tonsils coagulation | CIM and symptoms worsened, syrinx stable | unchanged |
| 44          | 5           | 7                    | normal                         | /             | none                             | CIM stable, symptoms improved | /                      |
| 45          | 17          | 14                   | normal                         | /             | bony PF decompression + C1 laminectomy | CIM and symptoms improved | /                      |
| 46          | 18          | 9                    | normal                         | /             | bony PF decompression + C1 laminectomy | CIM and symptoms regressed | /                      |
| 47          | 10          | 5                    | normal                         | sharp waves and spikes | none | CIM unchanged | unchanged |
| 48          | 6           | 15                   | cervico-thoracic syringomyelia | /             | PF decompression + duraplasty | CIM improved, syrinx and symptoms regressed | /                      |
| 49          | 6           | 9                    | normal                         | slow waves    | bony PF decompression + C1 laminectomy | CIM improved, symptoms stable | disappeared |
| 50          | 7           | 12                   | normal                         | /             | bony PF decompression + C1 laminectomy | CIM stable, symptoms improved | /                      |
| 51          | 12          | 28                   | holocord syringomyelia        | /             | PF decompression + duraplasty | CIM, syrinx and symptoms improved | /                      |
| 52          | 13          | 7                    | normal                         | /             | none                             | CIM unchanged            | /                      |
| 53          | 5           | 11                   | normal                         | /             | ICP monitoring                   | CIM unchanged            | /                      |
| 54          | 5           | 11                   | cervico-thoracic syringomyelia | /             | PF decompression + duraplasty | CIM and symptoms improved, syrinx unchanged | /                      |
| 55          | 8           | 9                    | normal                         | /             | bony PF decompression + C1 laminectomy | CIM stable, symptoms regressed | /                      |
| 56          | 2           | 17                   | normal                         | /             | bony PF decompression + C1 laminectomy | CIM and symptoms improved | /                      |
| 57          | 3           | 5                    | normal                         | /             | none                             | CIM unchanged            | /                      |
| 58          | 8           | 8                    | cervico-thoracic hydromyelia  | /             | bony PF decompression + C1 laminectomy | CIM and hydrocmyelia regressed, symptoms stable | /                      |
| 59          | 3           | 9                    | normal                         | /             | bony PF decompression + C1 laminectomy | CIM regressed, symptoms improved | /                      |
| 60          | 15          | 12                   | ventriculomegaly              | /             | ETV                              | CIM and ventriculomegaly improved | /                      |
| 61          | 10          | 10                   | normal                         | /             | bony PF decompression + C1 laminectomy | CIM and symptoms regressed | /                      |
| 62          | 8           | 10                   | normal                         | /             | none                             | CIM unchanged            | /                      |
| 63          | 7           | 31                   | cervical syringomyelia        | /             | PF decompression + duraplasty | CIM regressed, syrinx improved, symptoms stable | /                      |
### Table 4. Cont.

| Patients No | Age (Years) | Tonsils Ectopia (mm) | Spinal Cord and Other Findings | EEG Anomalies | Surgery | CIM/Syrinx Outcome | EEG Anomalies Outcome |
|-------------|-------------|----------------------|-------------------------------|---------------|---------|-------------------|----------------------|
| 64          | 6           | 9                    | thoracic syringomyelia        | /             | none    | CIM unchanged      | /                    |
| 65          | 14          | 9                    | thoracic syringomyelia        | /             | bony PF decompression + C1 laminectomy | CIM and symptoms regressed | / |
| 66          | 6           | 6                    | normal                        | /             | none    | CIM spontaneously regressed | / |
| 67          | 9           | 21                   | cervico-thoracic syringomyelia | /             | bony PF decompression + tonsils coagulation | CIM and syrinx regressed, symptoms improved | / |
| 68          | 6           | 12                   | normal                        | theta activity | bony PF decompression + C1 laminectomy | CIM improved, symptoms stable | unchanged |
| 69          | 15          | 18                   | cervico-thoracic syringomyelia | /             | PF decompression + tonsils coagulation | CIM and symptoms regressed, syrinx improved | / |
| 70          | 7           | 12                   | normal                        | /             | bony PF decompression + C1 laminectomy | CIM stable, symptoms improved | / |
| 71          | 16          | 18                   | normal                        | slow waves    | bony PF decompression + C1 laminectomy | CIM improved, symptoms regressed | disappeared |
| 72          | 5           | 13                   | normal                        | /             | bony PF decompression + C1 laminectomy | CIM improved, symptoms regressed | / |
| 73          | 12          | 15                   | cervico-thoracic hydromyelia  | /             | bony PF decompression + C1 laminectomy | CIM and symptoms improved, syrinx regressed | / |
| 74          | 9           | 8                    | ventriculomegaly              | /             | ETV     | CIM improved and ventriculomegaly regressed | / |
| 75          | 9           | 7                    | cervical syringomyelia        | /             | bony PF decompression + C1 laminectomy | CIM and symptoms stable | / |
| 76          | 7           | 11                   | normal                        | /             | bony PF decompression + C1 laminectomy | CIM regressed, symptoms improved | / |
| 77          | 9           | 22                   | cervical syringomyelia        | /             | PF decompression + duraplasty | CIM and syrinx regressed, symptoms improved | / |

PF: posterior fossa; ETV: endoscopic third ventriculostomy.

### 3.3. Comparison between the Two Groups

No statistically significant differences between Group 1 and Group 2 were detected regarding M/F ratio (0.93 vs. 1.24, respectively) and presence of syringomyelia/hydromyelia (27% vs. 39%). Moreover, no relevant differences were found in the CM1/syringomyelia outcome, namely, about regression/improvement of preoperative signs/symptoms (87.5% vs. 76%), radiological regression/improvement of tonsillar ectopia (88% vs. 76%), and syringomyelia/hydromyelia (87.5% vs. 91%). In addition, no correlation was found between seizure-free condition and PF decompression in Group 1, or between disappearance of EEG anomalies and PF decompression in Group 2, although the significance of these data is not reliable because the number of observations is too small (29 and 13 cases, respectively) (Table 5).

On the other hand, a significant difference between the two groups was noticed in terms of mean age at admission (6.2 vs. 8.9 years, \( p = 0.003 \)), amount of tonsillar herniation (7.5 vs. 11.8 mm, \( p < 0.00001 \)), and PF decompression (27.5% vs. 70%, \( p = 0.0001 \)). A final, obvious difference was found in the occurrence of EEG anomalies (100% vs. 17%, \( p < 0.00001 \)) (Table 5).
Table 5. Comparison between the two groups.

|                         | Group 1 | Group 2 | p-Value          |
|-------------------------|---------|---------|-----------------|
| No patients             | 29      | 77      | Not significant |
| Mean age at admission   | 6.2 years | 8.9 years | 0.003           |
| M/F ratio               | 0.93    | 1.26    | Not significant |
| Presence of seizures    | All cases | None    | Not calculated (patients’ selection) |
| EEG anomalies           | All cases | 17%    | <0.00001        |
| Presence of CM1/syrinx symptoms | 27%    | 75%    | Not calculated (patients’ selection) |
| Mean tonsillar herniation | 7.5 mm | 11.8 mm | <0.00001        |
| Syringomyelia/hydromyelia | 27%    | 39%    | Not significant |
| Ventriculomegaly        | 7%      | 4%      | Not significant |
| Surgery for CM1/syrinx  | 31%     | 76.5%   | 0.0001          |
| Seizure-free            | 62%     | /       | Not calculated (patients’ selection) |
| Clinical CM1/syrinx improvement | 6/9 cases | 41/54 cases | Not significant |
| Imaging CM1 improvement | 8/9 cases | 41/54 cases | Not significant |
| Imaging syrinx improvement | 7/8 cases | 21/23 cases | Not significant |

4. Discussion

Several studies, conducted mainly at the end of the last century and the beginning of the current one, raised the possible physiopathological association between CM1 and epilepsy. However, although this was clinically hypothesized, it has not been demonstrated yet [9–13]. The aim of the present study was to provide an update on possible clinical correlations between these two conditions, taking into account the evolution in the classification of both epilepsy and CM1. Indeed, the present study, if compared with the previous ones, included the new classification of epilepsy [7] and the new concepts relating to the CIM definition, which is no longer considered a mere tonsillar descent alone [3]. Moreover, the current study is based on a pediatric population admitted at a single institution and investigated homogeneously.

A first issue resulting from our study concerns the epidemiological implications of the CM1 and epilepsy association. Historical, large surgical or radiological series showed a prevalence of epilepsy among CM1 patients ranging from 3.8% to 10% [14–16]. In the present experience, 18% of 157 CM1 patients followed in the considered 5-year period (77 children symptomatic for CM1, which are here included, and 80 asymptomatic) were found to present epileptic seizures. The main explanation for this higher rate may be selection bias, because our institution is a tertiary referral center for both CM1 and epilepsy; therefore, the number of symptomatic subjects with both conditions is concentrated. However, in more recent studies, such as that of Marianayagam and coworkers, similar rates are reported (13.8%) [17]. This could depend on the increasing number of CM1 diagnoses and on the more extensive diagnostic work-up that epileptic (and CM1) patients currently undergo. Such an epidemiological association is also well-known in Cavalier King Charles Spaniels dogs (CM1 77%, epilepsy 28%) where, however, no relevant correlations between the conditions have been found [18]. Similar considerations can be made about the prevalence of CM1 in epileptic subjects, which was as high as 12% in the present series but about 5% in historical series [19]. On the other hand, the occurrence of ventriculomegaly or hydrocephalus does not seem to be specific to our series, since it was sporadic and did not affect the occurrence of seizures in both groups. The treatment of this condition was based
on ETV because it has been largely demonstrated that CM1-associated hydrocephalus is obstructive [20]. ETV was used as first-line treatment in the management of such an association, and also achieved good results for the improvement of CM1 or syringomyelia, as demonstrated elsewhere [21]. ETV results also a “protective” factor against epilepsy in hydrocephalic children if compared to a ventriculo-peritoneal shunt [22,23].

A second issue is related to the characteristics of patients with epilepsy and CM1. According to the literature, benign focal epileptiform discharges (mainly in the frontal and in the centrotemporal regions) and complex partial seizures are the most common types of seizures, followed by partial seizures with/without generalization, generalized tonic seizures, and myoclonic seizures [10–13,15,19]. Generally, a good epileptic outcome is reported. The present series partially confirms the trend of the types of epilepsy, with a prevalence of focal seizures (both FM and FNM, 48%) over generalized seizures (A, GT, GTC, M, 39%) and with a 13% rate of FGCT. Instead, the good epileptic outcome is totally confirmed, 62% of patients being seizure-free and 31% showing a >90% reduction in their seizures (see Table 3). The types of seizures and their outcome do not allow the formulation of relevant hypotheses but only the acknowledgement of the differences that exist among the different series. The type of seizures, in particular, depends on the characteristics of enrolled patients. Our series offers a reliable panorama of epilepsy in CM1 children because it is composed only of consecutive patients without possibly confounding factors (such as tumors or other lesions, syndromes etc.). On the other hand, the usually good epileptic outcome could be explained by the low rate of children with encephalopathy or severe brain damage, which are sporadically associated with CM1 (or which, sometimes, may “obscure” the presence of CM1, so that it could not be adequately reported in this subset of patients). This is quite different from the observations in general series of epileptic patients, where some factors, such as young age, generalized seizures, or use of a combination of drugs, can predict the remission of seizures [24].

The clinical characteristics of our series confirm another typical aspect of epileptic children with CM1, which is the high rate of associated developmental delay. In Group 1, indeed, cognitive and/or behavioral problems affected about 45% of cases (see Table 1) and, in the literature, this rate is even higher, reaching about 100% in some series [10–13,15,19]. This association is the most important clinical factor at the base of the theories postulating the presence of a common denominator between epilepsy and CM1 that can explain their possible intercorrelation. The brain (and cerebellar) microdysgenesis was hypothesized, in old studies, as a common denominator of a possible maldevelopmental process explaining the co-existence of CM1 and epilepsy. Grosso et al., for example, proposed the hypothesis that the association among CM1 and mental retardation, speech delay, and epilepsy (that they found in nine children) is not occasional but could be the expression of a specific developmental disorder [12]. However, such an association was purely clinical, and no genetic validation was provided. Moreover, not all the reported patients showed epilepsy and/or EEG anomalies. No significant correlation between tonsillar descent and epilepsy was found. Similarly, Brill et al. described a series of 11 CM1 children with epilepsy, developmental delay in motor or language function, and autistic features in some of them [10]. The authors formulated the hypothesis of a “subtle cerebral dysgenesis” that could give reason of a non-incidental occurrence of CM1 in this subset of patients. A comparison with Chiari II malformation (CM2) and its associated, dysgenetic brain findings was used to support this theory. However, CM2 is a completely different entity and, once again, the number of studied cases was too small, and no controls were analyzed. The hypothesis of a microdysgenesis was also promoted by Elia et al. on a similar series of seven children based on the absence of manifest dysgenetic findings [11]. An attempt at demonstrating the presence of a microdysgenesis was realized by Iannetti and coworkers by SPECT in four children with CM1 and epilepsy [13]. The authors found hypoperfusion brain areas in all cases, and cerebellar hypoperfusion areas in two of them. Unfortunately, such an interesting study was not replicated in a larger number of cases. Actually, no conclusions can be deduced from such a small number of cases, especially considering that the presence of cerebellar anomalies in only two cases could represent an
incidental association. Moreover, one could speculate that the cerebellar anomalies could be the result of an impairment on the connectivity between the brain cortex and cerebellum, as recently demonstrated [25]. However, in these instances, epilepsy is thought to cause cerebellar anomalies, while it would not result from them or, considering the topic of the present article, from CM1.

On these grounds, the aforementioned studies did not prove a relationship between CM1 and epilepsy even though the occurrence of a syndrome, which could support this hypothesis, cannot be ruled out. Actually, the presence of a syndrome might justify the co-existence of CM1 and epilepsy because the syndrome itself can encompass a spectrum of neurological disorders including these two entities. In syndromes, seizures usually result from the syndrome-associated lesions, as happens, for example, in neurofibromatosis-1 [26,27], or from the brain damages included in the syndrome, as occurs, for example, in fetal alcohol spectrum disorders [28]. An association between CM1 and epileptogenic anomalies (e.g. focal cortical heterotopia, cortical dysplasia, holoprosencephaly) is proved outside syndromes [29]. However, although it cannot be excluded, a shared genetic and embryologic maldevelopment pattern between these malformations and CM1 is not demonstrated yet [30]. KCTD7 deficiency is an example of a shared genetic pathway involving the brain (epilepsy) and cerebellum (degeneration of Purkinje cells) [31] and could be used as a possible model for research on this topic. To date, a syndrome with a “pure” association between CM1 and epilepsy has not been identified in spite of the recent description of new syndromes, such as the PTEN hamartoma tumor or the SETD2 neurodevelopmental one, which include CM1 and epilepsy [32,33]. Once again, it is worth noting that, also in these syndromes, seizures are likely to result from associated brain anomalies (brain tumors, vascular malformations, ventriculomegaly, macrocephaly) rather from cerebellar abnormalities or CM1. To date, there is no evidence that CM1 or syringomyelia can affect the electrical activity of the brain to cause seizures.

A further issue for discussion is the potential epileptogenicity of the cerebellum and, as a consequence, of CM1. Although it is a retrospective analysis, the present study did not highlight the cerebellar anomalies on neuroimaging (even in children undergoing perfusion sequences or 3T MRI) of both groups of patients, apart those related to CM1. Buoni et al. reported on three children with asymptomatic CM1 and EEG anomalies whose EEG abnormalities regressed after PF decompression [9]. The authors theorized that the CM1-dependent CSF flow reduction (found during surgery) could interfere with the brain electrical activity (rather than possible cerebellar anomalies), which is therefore normalized after the restoration of a proper CSF flow. Several limits (some of them acknowledged by the authors themselves) affect this study and its intriguing hypothesis: (1) the patient sample is too small; (2) EEG could have improved spontaneously over the time; (3) one out of three children was epileptic, thus showing different reasons than CM1 to show seizures; in this case, however, IRDA, spikes, and spikes waves disappeared after surgery, but it is not reported if the patient received antiepileptic drugs; (4) the interpretation of EEGs may have not been correct; and (5) a microdysgenesis was not demonstrated (although it cannot be denied). Several previous studies supported the hypothesis of the role of the cerebellum in generating epilepsy, often based on animal experimental models [10,28–30]. Nevertheless, experimental and clinical studies failed in demonstrating an effect of cerebellar stimulation in epilepsy [34,35]. The cerebellum could be the target for a stimulation in patients with epilepsy-induced degeneration of cortical-cerebellar pathways, which could improve the course of epilepsy [25]. Of course, this is something different from supporting a systematic role of cerebellum in epileptogenesis, which was sporadically demonstrated in anecdotal cases with cerebellar masses [36,37]. Recent volumetric studies comparing epileptic subjects and healthy controls, indeed, showed volumetric anomalies in the brain, lateral ventricles, and putamen, but not in the cerebellum [38]. On these grounds, the role of the cerebellum and, in particular, CM1, in epilepsy remains undemonstrated. This conclusion is further supported by some confusing symptoms mimicking seizures in CM1 patients (and, thus, potentially reducing the prevalence of “true” seizures in CM1), such as cerebellar fits, sleep disordered-breathing, paroxysmal dyskinesias, and syncope [15,39,40].
In summary, the clinical and experimental experiences from the literature lead to the conclusion that the association between epilepsy and CM1 is random [15,17]. The present study adds some clinical findings supporting the hypothesis of such a casual association. They can be summarized as follows: (1) epileptic patients (Group 1) show well-defined EEG patterns; in contrast, these are missing in CM1 patients, where EEG anomalies are not numerically relevant nor electrically specific but sporadic (Group 2); (2) no MRI anomalies suggesting microdysgenesis were found (in both Groups); (3) children with epilepsy and CM1 represent a substantially different population of patients compared with those with CM1 alone, as shown by some common, crucial findings, such as age at admission, amount of tonsillar descent, and need of PF decompression; and (4) the course and the outcome of one condition is not affected by the other, with the outcome of epilepsy depending on the drug treatment, while that of CM1/syringomyelia depending on surgery. These considerations were shared by most of the international experts attending the 2019 Consensus Conference on CM1 and Syringomyelia as part of their daily clinical experience [3]. Indeed, 94.1% of these experts concluded that, “in children with epilepsy and CM1, surgical treatment of CM1 does not improve the seizure disorder” because of the random association between the two conditions.

Limitations of the Study

A first limit of the present study is the retrospective analysis of a relatively small number of patients. Indeed, although it is the first study specifically addressing the problem of CM1 and epilepsy in the modern era, the small sample of patients does not allow the results to be considered as definitive. Moreover, as mentioned in the text, the statistical analysis was not reliable, in some instances, for the same reason. A multi-centric and prospective study would allow this limit to overcome to be overcome.

A second limit is the use of “rough” parameters for the quantification of CM1. We are planning to analyze new cases with multi-parameter and 3D measurements of the posterior fossa.

Author Contributions: Data curation, L.M., D.P., I.C., C.V., D.R.P.C. and R.C.; Formal analysis, L.M. and D.P.; Investigation, L.M., I.C., C.V., D.R.P.C. and R.C.; Methodology, L.M. and D.I.B.; Project administration, L.M.; Supervision, L.M. and G.T.; Validation, L.M.; Writing—original draft, L.M. and D.P.; Writing—review and editing, L.M., G.T. and D.I.B. All authors have read and agreed to the published version of the manuscript.

Funding: This research received no external funding.

Institutional Review Board Statement: The study was conducted in accordance with the Declaration of Helsinki and approved by the Institutional Ethics Committee of Fondazione Policlinico Universitario A. Gemelli Hospital, Rome (n. 4971, 12 April 2021).

Informed Consent Statement: Consent waived because of observational study.

Conflicts of Interest: The authors declare no conflict of interest.

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